

STIC Search Report Biotech-Chem Library

STIC Database Tracking Number

TO: Marcela Cordero Garcia

Location: 3c35/3c18

Art Unit: 1654

Wednesday, August 03, 2005

Case Serial Number: 10/722843

From: Noble Jarrell

Location: Biotech-Chem Library

Rem 1B71

Phone: 272-2556

Noble.jarrell@uspto.gov

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Other

Scientific and Technical Information Center

SEARCH REQUEST FORM

Act Unit. 1574 Phone Number: 2-2139 Results Format Preferred (circle): PAPER DIS Location (Biglogrount): Embeds (Mailton 9): 508 Results Format Preferred (circle): PAPER DIS To ensure an efficient and quality search, please attach a copy of the cover sheet, claims, and abstract or fill out the following: Title of Invention: PEPT (DES. INHICH TARGET Inventors (please provide full names): SET BOS Earliest Priority Date: 11/25/02 Search Taple: The experiment is detailed sustained of the search rapic, and describe as specifically as possible the subject matter to be searched. Inche describe species or structures, keywords, synonyms, evenyms, and registry numbers, and combine with the concept or utility of the inventional magning of the inventional magnine species in entire to the searched. Inche describe species or structures, keywords, synonyms, evenyms, and registry numbers, and combine with the concept or utility of the inventional magnines specied magnine. Public magnines specied magnine, or structures, keywords, synonyms, evenyms, and registry numbers, and combine with the concept or utility of the inventional magnines of the inventional magnines. **For Sequence Searches Only** Please include all pertinent information (purent, child, divisional, or issued patent numbers) along magnines are registrated to the inventional magnines. **PLEASE SEARCH BROAD CLM 6** **STAFF USE ONLY** **Type of Search **PLEASE SEARCH BROAD CLM 6** **PLEASE SEARCH BROAD CLM 6** **STAFF USE ONLY** **Type of Search **PLEASE SEARCH BROAD CLM 6** **PLEASE SEARCH BROAD CLM 6** **STAFF USE ONLY** **Type of Search **STAFF USE ONLY** **Type of Search **PLEASE SEARCH BROAD CLM 6** **STAFF USE ONLY** **STAFF USE ONLY** **Type of Search **STAFF USE ON	Requester's Full Nan	ne, MARCONA M	CORDERO GARCIA	Examine	er # : 80381	Date: 1/27/05
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Title of Invention: PEPT (DES. WHICH TARGET Inventors (please provide full names): SET BOS Earliest Priority Date: H/25/02 Search Topic: Please provide dull names): SET BOS Earliest Priority Date: H/25/02 Search Topic: Please provide detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. Included elected species or structures, heywords, symanyms, acromyms, and registry numbers, and combine with the concept or utility of the invencion of the search of the search of the invencence of the search of the invence of the search of the invencence of the invenc	Location (Bldg/Room#	(): REM3C3S (N	//ailbox #): <u>30/8</u> ************	. Kesults FC	e************	**********
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FILE 'HCAPLUS' ENTERED AT 08:23:29 ON 03 AUG 2005 3 US2005020810/PN OR (US2002-429174# OR US2003-475539#)/AP,PRN T.1

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FILE 'REGISTRY' ENTERED AT 08:24:34 ON 03 AUG 2005 209 SEA L2 L3

FILE 'WPIX' ENTERED AT 08:24:39 ON 03 AUG 2005 L4

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FILE COVERS 1907 - 3 Aug 2005 VOL 143 ISS 6 FILE LAST UPDATED: 2 Aug 2005 (20050802/ED)

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This file contains CAS Registry Numbers for easy and accurate substance identification.

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- ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN L1
- 2004:610128 HCAPLUS AN
- 141:157478 DN
- Entered STN: 30 Jul 2004 ED
- Peptides which target tumor and endothel al cells, compositions and uses
- Allan, Amy L.; Yoon, Won Hyung; Gladstone, Patricia L.; Ternansky, Robert IN J.; Parry, Graham; Donate, Fernando; Mazar, Andrew Attenuon, Llc, USA
- PA
- so PCT Int. Appl., 117 pp.

CODEN: PIXXD2

- DT Patent
- English LΑ
- IC ICM C07K
- 34-3 (Amino Acids, Peptides, and Proteins) Section cross-reference(s): 1, 63

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	WO 2004063213	A2	20040729	WO 2003-US37895	20031125 <
	WO 2004063213	A3	20050303		•

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W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
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     US 2004162239
     US 2005020810
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                                                                      20031125 <--
PRAI US 2002-429174P
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                                 20021125
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     US 2003-475539P
                                 20030602 <--
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CLASS
                 CLASS PATENT FAMILY CLASSIFICATION CODES
PATENT NO.
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                 ICM
 WO 2004063213
                         514/012.000; 514/013.000; 514/014.000; 514/015.000;
                 NCL
 US 2004162239
                         514/016.000; 514/017.000; 514/018.000; 530/324.000;
                         530/325.000; 530/326.000
                         530/324.000; 530/325.000; 530/326.000; 530/327.000;
US 2005020810
                 NCL
                         530/328.000; 530/329.000
OS
     MARPAT 141:157478
     The invention relates generally to peptide analogs of Ac-PHSCN-NH2 which
AΒ
     target tumor and endothelial cells and have antitumor, antiangiogenic and
     antimetastatic activity and to methods for their synthesis and use in
     pharmaceutical compns. for treating, preventing and detecting diseases
     characterized by tumor growth, metastasis and angiogenesis. The peptide
     analogs may serve, inter alia, as carriers of radioactivity, PET-active
     compds., toxins, fluorescent mols. and PEG mols. Peptides
     R1 [(NHCHR2CO) 0-1 (X1) 0-100] m-X2-X3-X4-X5-X6-[(X7) 0-1 (NHCHR3CO) 0-1] nNR4R5
     [R1 is (un) substituted acyl, alkyl, cycloalkyl or imino, or acyl chelate;
     R2 is substituted alkyl; R4, R5 are (un) substituted alkyl; X1, X7 are
     NH(CH:CH)1-6CO, NH(CH2)1-6CO, NHCHMeCO; X2-X6 are \alpha-amino acids
     which are defined; m, n are 0 or 1, with the proviso that R1 is not acetyl
     when R4 and R5 are H and m and n are 0] are claimed. Thus,
     Ac-Pro-His-Ser-Cys(Ac)-Asn-OH was prepared by the solid-phase method and
     coupled to doxorubicin hydrochloride to afford the conjugate.
     peptide prolylhistidylserylcysteinylaspartamide analog prepn antitumor
ST
     Angiogenesis
     Angiogenesis inhibitors
     Antitumor agents
     Neoplasm
        (preparation of peptides which target tumor and endothelial cells)
IT
     Peptides, preparation
     RL: DGN (Diagnostic use); PAC (Pharmacological activity); SPN (Synthetic
     preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (preparation of peptides which target tumor and endothelial cells)
     Polyoxyalkylenes, reactions
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (preparation of peptides which target tumor and endothelial cells)
IT
     729594-60-9P
     RL: DGN (Diagnostic use); PAC (Pharmacological activity); RCT (Reactant);
     SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological
     study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
        (preparation of peptides which target tumor and endothelial cells)
     7440-74-6DP, Indium, complexes with DPTA peptide conjugate
TT
                              729594-61-0P
                                              729594-62-1P
                                                               729594-63-2P
     262438-43-7DP, analogs
                                                                    729594-68-7P
                                                    729594-67-6P
                                    729594-66-5P
     729594-64-3P
                     729594-65-4P
                     729594-70-1P
                                     729594-71-2P
                                                    729594-72-3P
                                                                     729594-73-4P
     729594-69-8P
                                                    729594-77-8P
                                                                     729594-78-9P
     729594-74-5P
                     729594-75-6P
                                     729594-76-7P
                                                    729594-82-5P
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     729594-79-0P
                                     729594-81-4P
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                                                    729594-87-0P
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     729595-07-7P 729595-08-8P 729595-09-9P 729595-14-6P 730960-54-0P
     731003-01-3DP, Indium complexes 731003-01-3P 731003-02-4P
     RL: DGN (Diagnostic use); PAC (Pharmacological activity); SPN (Synthetic
     preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (preparation of peptides which target tumor and endothelial cells)
     456-22-4, 4 Fluorobenzoic acid 501-97-3 553-12-8 3301-79-9, 6
     Carboxyfluorescein 13811-11-5 25316-40-9, Doxorubicin hydrochloride
     34071-95-9 66134-67-6 76823-03-5, 5 Carboxyfluorescein 106966-68-1
     137076-54-1, 1,4,7,10-Tetraazacyclododecane-1,4,7,10-tetraacetic acid,
     tris 1 1 dimethylethyl ester 517913-89-2 622405-78-1 729595-15-7
     729595-16-8D, resin-bound 729595-17-9
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (preparation of peptides which target tumor and endothelial cells)
     729595-10-2DP, resin-bound 729595-11-3DP, resin-bound resin-bound 729595-13-5DP, resin-bound
ΙT
                                                                   729595-12-4DP,
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation of peptides which target tumor and endothelial cells)
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     ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN
     2004:467702 HCAPLUS
ΑN
     141:33798
DN
     Entered STN: 10 Jun 2004
ED
     Peptides which inhibit angiogenesis, cell migration, cell invasion and
TΙ
     cell proliferation, their preparation, and compositions and therapeutic
     uses thereof
     Allan, Amy L.; Donate, Fernando; Hopkins, Stephanie A.; Gladstone,
IN
     Patricia L.; Mazar, Andrew; O'Hare, Sean M.; Parry, Graham; Plunkett,
     Marian L.; Ternansky, Robert J.; Yoon, Won Hyung
     Attenuon, LLC, USA
PΑ
     PCT Int. Appl., 88 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
IC
     ICM A61K
     1-8 (Pharmacology)
     Section cross-reference(s): 34, 63
FAN.CNT 2
                                              APPLICATION NO.
                                                                      DATE
     PATENT NO.
                          KIND
                                  DATE
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                                              WO 2003-US38175
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              LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
         PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
              ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
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     US 2005020810
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PRAI US 2002-429174P
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     US 2003-475539P
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                  CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
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 WO 2004047771
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                  NCL .
                          514/016.000; 514/017.000; 514/018.000; 530/324.000;
                          530/325.000; 530/326.000
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US 2005020810
                 NCL
                        530/328.000; 530/329.000
     MARPAT 141:33798
os
     The invention discloses peptides which inhibit angiogenesis, cell
AB
     migration, cell invasion and cell proliferation, as well as methods of
     making the peptides, pharmaceutical compns. containing the peptides, and
     methods of using the peptides and pharmaceutical compns. to treat diseases
     associated with aberrant vascularization, e.g. cancer.
     peptide cell invasion migration proliferation inhibition; antitumor
ST
     aberrant vascularization disease peptide prepn
TТ
     Sarcoma
        (cartilage chondrosarcoma; peptide inhibitors of angiogenesis, cell
        migration, cell invasion and cell proliferation, preparation, and compns.
        and therapeutic uses)
     Cartilage, neoplasm
ΙT
        (chondrosarcoma; peptide inhibitors of angiogenesis, cell migration,
        cell invasion and cell proliferation, preparation, and compns. and
        therapeutic uses)
IT
     Intestine, neoplasm
        (colon; peptide inhibitors of angiogenesis, cell migration, cell
        invasion and cell proliferation, preparation, and compns. and therapeutic
        uses)
IT
     Blood vessel
        (endothelium; peptide inhibitors of angiogenesis, cell migration, cell
        invasion and cell proliferation, preparation, and compns. and therapeutic
ΙT
     Blood vessel, neoplasm
     Sarcoma
        (hemangiosarcoma; peptide inhibitors of angiogenesis, cell migration,
        cell invasion and cell proliferation, preparation, and compns. and
        therapeutic uses)
TТ
     Angiogenesis
     Angiogenesis inhibitors
     Antitumor agents
     Brain, neoplasm
     Drug delivery systems
     Kidney, neoplasm
     Mammary gland, neoplasm
     Neoplasm
     Prostate gland, neoplasm
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
TТ
     Endothelium
        (vascular; peptide inhibitors of angiogenesis, cell migration, cell
        invasion and cell proliferation, preparation, and compns. and therapeutic
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     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     THU (Therapeutic use); BIOL (Biological study); USES (Uses)
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        cell proliferation, preparation, and compns. and therapeutic uses)
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     (Preparation); RACT (Reactant or reagent); USES (Uses)
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
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RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
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        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
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     2002:849621 HCAPLUS
ΑN
     137:353056
DN
     Entered STN: 08 Nov 2002
ED
     Preparation of benzenesulfonylpiperazines as matrix metalloproteinase
ΤI
     Chung, Yong-Jun; Lee, Keyong-Ho; Kim, Youn-Chul; Park, Ho-Jin
IN
     Kolon Ind. Inc., S. Korea PCT Int. Appl., 71 pp.
PA
SO
     CODEN: PIXXD2
DT
     Patent
     English
LA
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     ICM C07D403-12
     28-16 (Heterocyclic Compounds (More Than One Hetero Atom))
CC
     Section cross-reference(s): 1
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                                                                     DATE
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CLASS
                CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
 WO 2002088115 ICM C07D403-12
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C07C311/19; C07C311/29; C07D241/04; C07D241/08;
WO 2002088115
                 ECLA
                        C07D243/08; C07D245/02; C07D403/12+241B+207
                        4C063/AA01; 4C063/BB03; 4C063/BB08; 4C063/CC34;
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                         4C063/DD04; 4C063/DD12; 4C063/EE01; 4C086/AA01;
                        4C086/AA02; 4C086/AA03; 4C086/AA04; 4C086/BC49; 4C086/BC73; 4C086/GA07; 4C086/GA08; 4C086/GA09;
                         4C086/GA12; 4C086/MA01; 4C086/MA04; 4C086/NA14;
                         4C086/ZA33; 4C086/ZA44; 4C086/ZA45; 4C086/ZA67;
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                         4C086/ZB11; 4C086/ZB15; 4C086/ZB26; 4C086/ZC06;
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                         514/218.000; 514/254.010; 514/255.020; 514/183.000;
                 NCL
US 2004138206
                         540/575.000; 540/474.000; 544/372.000; 544/383.000
                         C07C311/19; C07C311/29; C07D241/04; C07D241/08;
                 ECLA
                         C07D243/08; C07D245/02; C07D403/12+241B+207
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MARPAT 137:353056 OS

GΙ

Title compds. [I; n = 0-3; A = CO2H, CONHOH, CH2SH, CH2OH; B = H, alkyl, NO2, aryl, heteroaryl, pyrrolyl, halo, alkoxy, aryloxy, alkylamino, alkylthio, CONHR, NHCOR, NHCO2R, NHCONHR, etc.; R = H, alkyl, aryl, heteroaryl, tetragonal to octagonal cyclic compound, alkyl substituted by a tetragonal to octagonal (hetero)cyclic compound; Z = H, O, S, provided that when Z = O, S it takes a double bond; Y = H, alkyl, aryl, heteroaryl, alkyl substituted by a tetragonal to octagonal cyclic compound, alkyl substituted by a tetragonal to octagonal heterocyclyl, CONHR, NHCOR, NHCO2R, NHCONHR, alkyl having a double or triple bond], were prepared Thus, Me 1-(4-methoxybenzenesulfonyl)-5-oxopiperazine-2-carboxylate (preparation given) was stirred 5 h with aqueous NH2OH to give 45% 1-(4methoxybenzenesulfonyl)-5-oxopiperazine-2-hydroxamic acid. This inhibited MMP-2 with IC50 = 0.004 μM . I are angiogenesis controlling materials that can inhibit overexpression of matrix metalloproteinase that decomps. protein constituents in extracellular matrix and basement membranes of connective tissues.

benzenesulfonylpiperazine prepn matrix metalloproteinase inhibitor; cancer stangiogenesis inhibitor prepn benzenesulfonylpiperazine; hydroxamate benzenesulfonylpiperazine prepn anticancer; piperazinehydroxamate arylsulfonyl prepn mmp inhibitor

IT Antitumor agents

Human

(preparation of benzenesulfonylpiperazines as matrix metalloproteinase inhibitors)

Hydroxamic acids ΤТ

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

(preparation of benzenesulfonylpiperazines as matrix metalloproteinase inhibitors)

TТ Angiogenesis Neoplasm

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IT
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     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
        (preparation of benzenesulfonylpiperazines as matrix metalloproteinase
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              Methylamine, reactions 98-68-0, 4-Methoxybenzenesulfonyl 100-46-9, Benzylamine, reactions 105-36-2, Ethyl bromo
     74-89-5, Methylamine, reactions
IT
                                                   105-36-2, Ethyl bromoacetate
     109-73-9, n-Butylamine, reactions 111-26-2, Hexylamine 111-86-4,
     Octvlamine 112-90-3, Oleylamine
                                         507-09-5, Thiolacetic acid, reactions
     696-59-3, 2,5-Dimethoxytetrahydrofuran 765-30-0, Cyclopropylamine
     2016-57-1, Decylamine 2038-03-1, N-(2-Aminoethyl)morpholine
                                                                     2706-56-1,
     2-(2-Aminoethyl)pyridine 3731-51-9, 2-Aminomethylpyridine
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     DL-Serine methyl ester hydrochloride
     27578-60-5, 1-(2-Aminoethyl)piperidine 202752-04-3
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     (Reactant or reagent)
        (preparation of benzenesulfonylpiperazines as matrix metalloproteinase
        inhibitors)
              THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
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RE
(1) Agouron Pharmaceuticals Inc; US 5753653 1996 HCAPLUS
(2) Anon; J MED CHEM 2000, V43(3), P369
(3) Fujisawa Pharmaceutical Co Ltd; WO 9827069 A 1998 HCAPLUS
(4) Nippon Soda Co Ltd; WO 0102371 A 2001 HCAPLUS
(5) Pfizer Inc; WO 9633172 A 1996 HCAPLUS
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    DOCUMENTATION NOW AVAILABLE IN DERWENT WORLD PATENTS INDEX
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    PLEASE CHECK:
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AN
     2004-561873 [54]
     2004-450190 [42]
CR
DNC
     C2004-205382
     New peptide derivatives having anti-tumor activity useful for the
TI
     treatment, prevention or detection of cancer.
DC
     ALLAN, A L; DONATE, F; GLADSTONE, P L; MAZAR, A; PARRY, G; TERNANSKY, R J;
IN
     YOON, W H
     (ATTE-N) ATTENUON LLC; (ALLA-I) ALLAN A L; (DONA-I) DONATE F; (GLAD-I)
     GLADSTONE P L; (MAZA-I) MAZAR A; (PARR-I) PARRY G; (TERN-I) TERNANSKY R J;
     (YOON-I) YOON W H
CYC
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                    A1 20050127 (200509)
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PRAI US 2003-475539P
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TC
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     ICS C07K007-06
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AB
     NOVELTY - Peptide derivatives (I) and their salts, solvates, hydrates or
     N-oxides are new.
          DETAILED DESCRIPTION - Peptide derivatives of formula (I) and their
    salts, solvates, hydrates or N-oxides are new.
     j, k = 0-1;
     p, q = 0-100;
          = 0-1;
          R1 = (substituted) acyl, acyl chelate, (substituted) alkyl,
     (substituted) cycloalkyl or (substituted) imino;
          R2 = 1-6C alkyl with at least H replaced by a substituents of NR6R7,
     -OR8, -CO2R9, -S(O)2R10, -P(OR11)OR12 or (substituted) aryl;
          R6-R12
                 = H or R1;
          X1 = NH(C=C)gCO-, NH(CH2)hCO- or NHCH(CH3)CO-;
          = 1-6:
          X2 = cyclic derivative of formula (i-iii);
          X3 = imidazole derivative of formula (iv);
          X4 = alcohol derivative of formula (v-vi);
       = 1-4;
          X5 = sulfonyl derivative of formula (vii);
```

```
R13 = H, (substituted) alkyl, (substituted) acyl, (substituted)
     arylalkyl, (substituted) aryl or -S(O)nR14;
       = 1-5;
          R14 = (substituted) alkyl, (substituted) acyl, (substituted)
     arylalkyl or (substituted) aryl;
    x, y = 0-2;
          X6 = amide derivative formula (viii);
       = 1-4;
         X7 = NH(C=C)dCO-, -NH(CH2)eCO or -NHCH(CH3)CO-;
          R3 = 1-6C alkyl with at least H replace by a substituent of
     -NR15R16, -OR17, -CO2R18, -S(O)nR19, -P(OR20)OR21 or (substituted) aryl;
          R4, R5 = H or (substituted alkyl); and
          R15-R21 = H, (substituted) acyl, acyl chelate, (substituted) alkyl,
     (substituted) cycloalkyl or (substituted) imino.
          Provided that R1 is not acetyl when R4 and R5 are H and r and s 0.
          ACTIVITY - Cytostatic; Antiangiogenic.
          Tests details are described but no results given.
          MECHANISM OF ACTION - None given
          USE - (I) are useful for the treament, prevention or detection of
     cancer (claimed), tumor growth, metastasis and angiogenesis.
     Dwg.0/0
FS
     CPI
FΑ
     AB; GI; DCN
     CPI: B02-D; B04-C01B; B04-C01C; B04-C01D; B04-C01E; B04-C01F; B04-C01G;
MC
          B04-N04A; B14-H01
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AN
     2004-561873 [54]
CR
DNC
     C2004-168702
     Novel peptides useful as e.g. angiogenesis inhibitors for treating or
TТ
     preventing cancer, e.g. breast cancer, renal cancer, brain cancer, colon
     cancer.
DC
     B03
     ALLAN, A L; DONATE, F; GLADSTONE, P L; HOPKINS, S A; MAZAR, A; O'HARE, S
IN
     M; PARRY, G; PLUNKETT, M; TERNANSKY, R J; YOON, W H; PLUNKETT, M L
     (ALLA-I) ALLAN A L; (DONA-I) DONATE F; (GLAD-I) GLADSTONE P L; (HOPK-I)
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     HOPKINS S A; (MAZA-I) MAZAR A; (OHAR-I) O'HARE S M; (PARR-I) PARRY G;
     (PLUN-I) PLUNKETT M; (TERN-I) TERNANSKY R J; (YOON-I) YOON W H; (ATTE-N)
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            YU ZA ZM ZW
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     US 2004162239
                    A1 20040618 (200471)
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     2003-475539P 20030602, US 2003-723144 20031125; AU 2003297609 A1 AU
     2003-297609 20031125
FDT AU 2003297609 Al Based on WO 2004047771
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IC
     ICS A61K038-10; C07K007-06; C07K007-08
     WO2004047771 A UPAB: 20041104
     NOVELTY - Peptides are new.
          DETAILED DESCRIPTION - Peptides of formula R1-Ax-By-C'z-(N-CH(R2)-
     C(O))a-(N-CH(R3)-C(O))b-R4 (I), their salt, solvates, hydrates or N-oxides
     are new.
```

```
a, b and x - z = 0 or 1;
     A = cyclic amino acid;
     B = basic amino acid;
     C' = small amino acid;
     R1 = (hetero)alkyl, acyl, alkylsulfonyl, (hetero)arylalkyl,
(hetero)arylsulfonyl or oxycarbonyl (all optionally substituted);
     R2 = alkyl, -(CH2)mS(0)nR5, -(CH2)mS(0)n-S(0)oR5 or -(CMe)mS(0)nR5;
m = 1-4;
n and o = 0-2;
     R3 = -CH2CONH2 \text{ or } -CH2CH2CONH2;
     R4 = alkyl, -NR6R7 \text{ or } -OR8;
     R5 = (hetero)alkyl, acyl, (hetero)aryl, (hetero)arylalkyl or
oxycarbonyl (all optionally substituted);
     R6, R7 = H or alkyl;
     R8 = (hetero)alkyl, (hetero)aryl or (hetero)arylalkyl(all optionally
substituted).
Provided that:
     (1) when m is 1, R5 is other than methyl;
```

- (2) a is 1 unless A is proline, B is histidine, C is serine;
- (3) when a is 0, b is 0; and
- (4) R2 is -(CH2)mS(O)nR5 or -(CH2)mS(O)n-S(O)oR5 unless b, x, y and z are 1.

An INDEPENDENT CLAIM is also included for treatment or prevention of cancer involving administering (I) optionally with an anti-cancer agent.

ACTIVITY - Cytostatic; Antiangiogenic; Antiarthritic; Antidiabetic; Antiarteriosclerotic; Ophthalmological; Vulnerary; Antirheumatic; Dermatological; Antipsoriatic; Antiparasitic; Osteopathic; Vasotropic; Tranquilizer; Thrombolytic; Gynecological; Antiinflammatory; Respiratory-Gen.; Antiulcer; Antisickling.

MECHANISM OF ACTION - Angiogenesis inhibitor; Cell migration, cell invasion and cell proliferation inhibitor; Tumor growth inhibitor.

Acetyl-Pro-His-Ser-Cys(S-tert-Bu)-Asn-NH2 (A) was tested in vivo for its ability to inhibit FGF-2 mediated angiogenesis in a Matrigel Plug (RTM) model according to Passaniti et al., 1992, Lab Invest. 67:519-528.

(A) showed % inhibition of 88.2 plus or minus 42.9.

USE - (I) Are used for treating or preventing cancer e.g. breast cancer, renal cancer, brain cancer, colon cancer, prostrate cancer, chondrosarcoma or angiosarcoma (claimed); for treating diseases associated with aberrant vascularization including arthritis, diabetes, arteriosclerosis, arteriovenous malformation, corneal graft neovascularization, delayed wound healing, diabetic retinopathy, age related macular degeneration, granulation burn, hemophilic joint, rheumatoid arthritis, hypertrophic scar, neovascular glaucoma, nonunion fracture, Osier Weber Syndrome, psoriasis, retrolental fibroplasia, pterygium, scleroderma, trachoma, vascular adhesion, ocular neovascularization, parasitic disease, hypertrophy following surgery, inhibition of hair growth, macular degeneration, osteoarthritis, benign hyperplasia, atherosclerosis, myocardial angiogenesis, post-balloon angioplasty vascular restenosis, neointima formation following vascular trauma, vascular graft restenosis, coronary collateral formation, deep venous thrombosis, ischemic limb angiogenesis; telangiectasia, pyogenic granuloma, corneal disease, rubeosis, neovascular glaucoma, diabetic and other retinopathy, retrolental fibroplasias, diabetic neovascularization, endometriosis, fibrosis associated with a chronic inflammatory condition, traumatic spinal cord injury including ischemia, scarring or fibrosis, lung fibrosis, chemotherapy-induced fibrosis; wound healing with scarring and fibrosis, peptic ulcers, a bone fracture, keloids, or a disorder of vasculogenesis, hematopoiesis, ovulation, menstruation, pregnancy or placentation associated with pathogenic cell invasion or with angiogenesis, retinopathy of prematurity, sickle cell retinopathy or retinal vein occlusion; for treating uterine disease; to detect or image disease or conditions associated with undesired cell migration, invasion or proliferation.

ADVANTAGE - The compounds (I) are potent inhibitors of angiogenesis, cell migration, cell invasion and cell proliferation.

Dwg.0/5

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FS
     CPI
     AB; DCN
ĖΑ
     CPI: B04-C01A; B06-H; B07-H; B10-A04; B10-A08; B10-A10; B10-A12C; B10-B02;
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          B14-N01; B14-N03; B14-N14; B14-N16; B14-N17; B14-P02; B14-R02;
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     ANSWER 3 OF 3 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN
L4
AN
     2003-103447 [09]
                       WPIX
DNC C2003-026138
ΤТ
     New sulfonamide derivatives useful in the treatment of e.g. cancer.
DC
     CHUNG, Y; KIM, Y; LEE, K; PARK, H; JUNG, Y J; KIM, Y C; LEE, G H; PARK, H
IN
     J; CHUNG, Y J
      (KOLO-N) KOLON IND INC; (CHUN-I) CHUNG Y; (KIMY-I) KIM Y; (LEEK-I) LEE K;
      (PARK-I) PARK H
CYC
     101
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                     A1 20021107 (200309)* EN
                                                 71
                                                       C07D403-12
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             LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT RO
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     20010426; KR 2003047127 A KR 2001-77522 20011207; KR 2003075322 A KR
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      20020424; AU 2002251588 A1 AU 2002-251588 20020424; US 2004138206 A1 WO
      2002-KR759 20020424, US 2003-475539 20031211; KR 432928 B KR
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PRAI KR 2002-14481
     KR 2001-77522
                           20011207
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     WO 200288115 A UPAB: 20030206
AΒ
      NOVELTY - New sulfonamide derivatives of formula (I), their optical
      isomers, salts or solvates.
          DETAILED DESCRIPTION - Sulfonamide derivatives of formula (I), their
      optical isomers, salts or solvates are new.
        = 0 -3;
           A = CO2H, CONHOH, CH2SH or CH2OH;
           B = H, 1-8C lower alkyl, nitro, aryl, heteroaryl, pyrrole, halo,
      1-8C O-lower alkyl, O-aryl, N-lower alkyl, S-lower alkyl, phenyl
      (substituted by X), amide compound of formula CONHR or NHCOR, carbamate
      compound of formula NHCOOR or urea compound of formula NHCONHR;
           X = H, 1-8C lower alkyl, 9-20C higher alkyl, 9-20C higher alkyl
      comprising a double bond, (hetero)aryl, halo, O-lower alkyl, O-aryl,
```

O-heteroaryl, N-aryl, N-heteroaryl, S-aryl, S-heteroaryl, 1-20C alkyl-amine derivative, 1-20C alkyl-carboxylic acid derivative, amine or nitro:

R = H, 1-8C lower alkyl, (hetero)aryl, tetragonal to octagonal (hetero)cyclic compound or 1-8C lower alkyl (substituted by tetragonal to octagonal (hetero)cyclic compound); Z = H, O or S;

Y = H, 1-18C alkyl, (hetero)aryl, 1-8C lower alkyl (substituted by a tetragonal to octagonal (hetero)cyclic compound), amide compound of formula CONHR or NHCOR, carbamate compound of formula NHCOOR, urea compound of formula NHCONHR, 1-8C lower alkyl having a double or a triple bond, 9-20C higher alkyl having a double or a triple bond.

Provided that when Z is O or S the C(ring atom)-Z bond is a double bond.

INDEPENDENT CLAIMS are also included for:

- Preparation of (I);
- (2) New 4-phenylsulfonyl-piperazine intermediates (II);
- (3) Preparation of (II) comprising reaction of a substituted phenylsulfamide of formula (III) with methanesulfonyl chloride, toluenesulfonyl chloride or triflic anhydride in the presence of a base, and reaction of the product with primary amine;
 - (4) New substituted phenylsulfamide of formula (III); and
- (5) Preparation of (III) comprising reaction of the compound of formula (IV) with ethyl bromoacetate and halogen in presence of an inorganic base and N,N-dimethyl formamide or acetonitrile solvent.

W and X = H, methyl, ethyl, t-butyl or 1-8C lower alkyl group comprising a benzyl group.

ACTIVITY - Cytostatic; Antiarteriosclerotic; Ophthalmological; Antidiabetic; Antiarthritic; Antirheumatic; Antiinflammatory; Antiulcer; Osteopathic; Antiseborrheic; Dermatological; Anti-HIV; Antipsoriatic; Vulnerary.

MECHANISM OF ACTION - Matrix metalloproteinase (MMP) inhibitor. The MMP inhibitor activities were measured by fluorescence assay as described by Knight, C. G., Willenbrock, F., Murphy, G. A., FEBS Lett. 1992, 296, 263-266. For 1-(4'-bromo-biphenyl-4-sulfonyl)-4-octyl-5-oxo-piperazine-2-hydroxamate. The results indicated an IC50 (mu M) valve of 0.016, 0.002, 0.0013 and 0.007 for MMP-1, MMP-2, MMP-9 and MMP-13 respectively.

USE - In the treatment of cancer metastasis, solid cancer and angiogenesis (claimed). Also useful in the treatment of cardiovascular disease (e.g. hemangioma, angiofibroma), angiostenosis, edematous sclerosis, eye diseases caused by angiogenesis, corneal transplantation, angiogenic glaucoma, diabetic retinopathy, angiogenic corneal disease, age-related macular degeneration, pterygium, retinal degeneration, retreolental fibroplasias, granular conjunctivitis, skin diseases caused by angiogenesis (e.g. chronic inflammatory diseases e.g. arthritis, psoriasis, telangiectasis, granuloma pyogenicum, sebborhoeic dermatitis), periodontal disease, tumors, rheumatoid arthritis, inflammation, hyperparathyroidism, diabetes, corneal ulcers, osteoporosis, stomach ulcers, wounds, wrinkles, acne, AIDS, burns, arteriosclerosis, bone fractures.

ADVANTAGE - The compound is a potent proteinase inhibitor. $\ensuremath{\text{Dwg.0/0}}$

FS CPI

FA AB; GI; DCN

MC CPI: B07-D03; B07-D11; B10-A08; B14-C03; B14-C09; B14-D07C; B14-E08; B14-F01; B14-F02F2; B14-F07; B14-G01B; B14-H01; B14-N01; B14-N03; B14-N06B; B14-N11; B14-N17; B14-S04; N02-F01

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=> d his full

1.2

L3

L5

L6

L19

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U OR "TERNANSKY ROBERT J"/AU OR "TERNANSKY ROBERT JOHN"/AU OR

"TERNANSKY ROBERTJ"/AU)

E TERNANSKY B/AU E ALLAN A/AU

L10 35 SEA ABBEON PLU=ON ("ALLAN A"/AU OR "ALLAN A L"/AU) E ALLAN AMY/AU

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E GLADSTONE P/AU

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L"/AU OR "GLADSTONE PATRICIA LOUISE"/AU OR "GLADSTONE PATRICIAL
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E YOON W/AU

L13 13 SEA ABB=ON PLU=ON ("YOON W"/AU OR "YOON W H"/AU)
E YOON WON/AU

L14 10 SEA ABBEON PLU=ON ("YOON WON"/AU OR "YOON WON HYUNG"/AU)
E YOON WONHY/AU
E PARRY G/AU

L15

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E PARRY GRAHAM/AU

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E DONATE F/AU

L17 25 SEA ABB=ON PLU=ON ("DONATE F"/AU OR "DONATE F A"/AU OR "DONATE F ERNANDO"/AU)

E MAZAR A/AU

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L29
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L33
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L34
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L35
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L37
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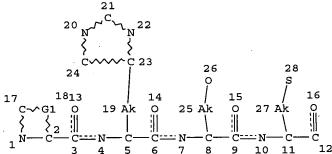
*
The CA roles and document type information have been removed from *
the IDE default display format and the ED field has been added, *
effective March 20, 2005. A new display format, IDERL, is now *
available and contains the CA role and document type information. *

*

Structure search iteration limits have been increased. See HELP SLIMITS for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

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DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 28

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100.0% PROCESSED 95013 ITERATIONS SEARCH TIME: 00.00.07

125 ANSWERS

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RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG US 2003-723144 US 2004162239 A1 20040819 20031125 20050127 US 2003-722843 20031125 US 2005020810 A1 PRAI US 2002-429174P Р 20021125 US 2003-475539P Р 20030602 CLASS CLASS PATENT FAMILY CLASSIFICATION CODES PATENT NO. ----______ WO 2004063213 CO7K ICM 514/012.000; 514/013.000; 514/014.000; 514/015.000; US 2004162239 NCL 514/016.000; 514/017.000; 514/018.000; 530/324.000; 530/325.000; 530/326.000 530/324.000; 530/325.000; 530/326.000; 530/327.000; US 2005020810 530/328.000; 530/329.000 os MARPAT 141:157478 The invention relates generally to peptide analogs of Ac-PHSCN-NH2 which target tumor and endothelial cells and have antitumor, antiangiogenic and

antimetastatic activity and to methods for their synthesis and use in

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pharmaceutical compns. for treating, preventing and detecting diseases
     characterized by tumor growth, metastasis and angiogenesis. The peptide
     analogs may serve, inter alia, as carriers of radioactivity, PET-active
     compds., toxins, fluorescent mols. and PEG mols. Peptides
     R1 [ (NHCHR2CO) 0-1 (X1) 0-100] m-X2-X3-X4-X5-X6- [ (X7) 0-1 (NHCHR3CO) 0-1] nNR4R5
     [R1 is (un) substituted acyl, alkyl, cycloalkyl or imino, or acyl chelate; R2 is substituted alkyl; R4, R5 are (un) substituted alkyl; X1, X7 are
     NH(CH:CH)1-6CO, NH(CH2)1-6CO, NHCHMeCO; X2-X6 are \alpha-amino acids
     which are defined; m, n are 0 or 1, with the proviso that R1 is not acetyl
     when R4 and R5 are H and m and n are 0] are claimed. Thus,
     Ac-Pro-His-Ser-Cys(Ac)-Asn-OH was prepared by the solid-phase method and
     coupled to doxorubicin hydrochloride to afford the conjugate.
     peptide prolylhistidylserylcysteinylaspartamide analog prepn antitumor
IT
     Angiogenesis
     Angiogenesis inhibitors
     Antitumor agents
     Neoplasm
        (preparation of peptides which target tumor and endothelial cells)
IT
     Peptides, preparation
     RL: DGN (Diagnostic use); PAC (Pharmacological activity); SPN (Synthetic
     preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (preparation of peptides which target tumor and endothelial cells)
IT
     Polyoxyalkylenes, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (preparation of peptides which target tumor and endothelial cells)
     729594-60-9P
     RL: DGN (Diagnostic use); PAC (Pharmacological activity); RCT (Reactant);
     SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological
     study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
        (preparation of peptides which target tumor and endothelial cells)
     7440-74-6DP, Indium, complexes with DPTA peptide conjugate
IT
     262438-43-7DP, analogs 729594-61-0P 729594-62-1P
     729594-63-2P 729594-64-3P 729594-65-4P
     729594-66-5P 729594-67-6P 729594-68-7P
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     729594-99-4P 729595-00-0P 729595-01-1P
                    729595-03-3DP, polyethylene glycol derivative
     729595-02-2P
     729595-04-4P 729595-05-5P 729595-06-6P
     729595-07-7P 729595-08-8P 729595-09-9P
     729595-14-6P 730960-54-0P 731003-01-3DP,
     Indium complexes 731003-01-3P 731003-02-4P
RL: DGN (Diagnostic use); PAC (Pharmacological activity); SPN (Synthetic
     preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (preparation of peptides which target tumor and endothelial cells)
     456-22-4, 4 Fluorobenzoic acid 501-97-3 553-12-8
                                                             3301-79-9, 6
TT
                                       25316-40-9, Doxorubicin hydrochloride
     Carboxyfluorescein 13811-11-5
                  66134-67-6 76823-03-5, 5 Carboxyfluorescein
                                                                   106966-68-1
     34071-95-9
     137076-54-1, 1,4,7,10-Tetraazacyclododecane-1,4,7,10-tetraacetic acid,
                                                   622405-78-1
                                                                  729595-15-7
     tris 1 1 dimethylethyl ester
                                     517913-89-2
     729595-16-8D, resin-bound 729595-17-9
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (preparation of peptides which target tumor and endothelial cells)
     729595-10-2DP, resin-bound
                                   729595-11-3DP, resin-bound
IT
                   729595-13-5DP, resin-bound
     resin-bound
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
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Absolute stereochemistry.

PAGE 1-B

English

LΑ

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L22 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN
AN
     2004:467702 HCAPLUS
DN
     141:33798
     Entered STN: 10 Jun 2004
ED
     Peptides which inhibit angiogenesis, cell migration, cell invasion and
ΤI
     cell proliferation, their preparation, and compositions and therapeutic
     uses thereof
IN
     Allan, Amy L.; Donate, Fernando; Hopkins, Stephanie
     A.; Gladstone, Patricia L.; Mazar, Andrew; O'Hare,
     Sean M.; Parry, Graham; Plunkett, Marian L.; Ternansky,
     Robert J.; Yoon, Won Hyung
PA
     Attenuon, LLC, USA
     PCT Int. Appl., 88 pp.
so
     CODEN: PIXXD2
DT
     Patent
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IC
     ICM A61K
CC
     1-8 (Pharmacology)
     Section cross-reference(s): 34, 63
FAN.CNT 2
                                  DATE
                                               APPLICATION NO.
                                                                        DATE
     PATENT NO.
                          KIND
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                                 20040610 WO 2003-US38175
                           A2
                                                                       20031125
ΡI
     WO 2004047771
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
             ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD,
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                                  20030602
     US 2003-475539P
CLASS
                         PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                  CLASS
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 WO 2004047771
                  ICM
                         A61K
                         514/012.000; 514/013.000; 514/014.000; 514/015.000;
 US 2004162239
                  NCL
                          514/016.000; 514/017.000; 514/018.000; 530/324.000;
                          530/325.000; 530/326.000
                         530/324.000; 530/325.000; 530/326.000; 530/327.000; 530/328.000; 530/329.000
 US 2005020810
                  NCL
os
     MARPAT 141:33798
     The invention discloses peptides which inhibit angiogenesis, cell
AB
     migration, cell invasion and cell proliferation, as well as methods of
     making the peptides, pharmaceutical compns. containing the peptides, and
     methods of using the peptides and pharmaceutical compns. to treat diseases
     associated with aberrant vascularization, e.g. cancer.
     peptide cell invasion migration proliferation inhibition; antitumor
ST
     aberrant vascularization disease peptide prepn
IT
     Sarcoma
         (cartilage chondrosarcoma; peptide inhibitors of angiogenesis, cell
        migration, cell invasion and cell proliferation, preparation, and compns.
        and therapeutic uses)
TT
     Cartilage, neoplasm
         (chondrosarcoma; peptide inhibitors of angiogenesis, cell migration,
        cell invasion and cell proliferation, preparation, and compns. and
        therapeutic uses)
IT
     Intestine, neoplasm
        (colon; peptide inhibitors of angiogenesis, cell migration, cell
        invasion and cell proliferation, preparation, and compns. and therapeutic
        uses)
     Blood vessel
IT
         (endothelium; peptide inhibitors of angiogenesis, cell migration, cell
        invasion and cell proliferation, preparation, and compns. and therapeutic
        uses)
TT
     Blood vessel, neoplasm
     Sarcoma
         (hemangiosarcoma; peptide inhibitors of angiogenesis, cell migration,
        cell invasion and cell proliferation, preparation, and compns. and
        therapeutic uses)
IT
     Angiogenesis
     Angiogenesis inhibitors
     Antitumor agents
     Brain, neoplasm
     Drug delivery systems
     Kidney, neoplasm
     Mammary gland, neoplasm
```

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Neoplasm
    Prostate gland, neoplasm
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
TТ
    Endothelium
        (vascular; peptide inhibitors of angiogenesis, cell migration, cell
        invasion and cell proliferation, preparation, and compns. and therapeutic
        uses)
IT
     701201-26-5D, biotinylated
    RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
    THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
    701200-82-0P 701201-01-6P
IT
    RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic
    preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); RACT (Reactant or reagent); USES (Uses)
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        cell proliferation, preparation, and compns. and therapeutic uses)
     81658-55-1P 701200-81-9P 701200-83-1P
TT
                    701200-85-3P 701200-86-4P
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     701200-84-2P
                   701200-89-7P 701200-90-0P
     701200-88-6P
     701200-91-1P 701200-92-2P 701200-93-3P
                                                  701200-97-7P
     701200-94-4P
                   701200-95-5P
                                  701200-96-6P
                                                                 701200-98-8P
     701200-99-9P 701201-00-5P 701201-02-7P
     701201-03-8P 701201-04-9P 701201-05-0P
     701201-06-1P 701201-07-2P 701201-08-3P
     701201-09-4P 701201-10-7P 701201-11-8P
     701201-12-9P 701201-13-0P 701201-14-1P
     701201-15-2P 701201-16-3P 701201-17-4P
     701201-18-5P 701201-19-6P 701201-20-9P
     701201-21-0P
                    701201-22-1P
                                   701201-23-2P 701201-24-3P
     701201-25-4P
     RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
     701201-28-7
TT
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
     98-88-4, Benzoyl chloride 100-39-0, Benzyl bromide 106-95-6, Allyl
TT
                        930-69-8
                                   1212-08-4, S-Phenyl benzenethiosulfonate
     bromide, reactions
     2719-27-9, Cyclohexanoyl chloride 2937-50-0, Allyl chloroformate
     2949-92-0, S-Methyl methanethiosulfonate 3282-30-2, Pivaloyl chloride
     5271-67-0, 2-Thiophenecarbonyl chloride 6482-24-2, 2-Bromoethyl
     methylether 7031-27-8, (Phenylthio) acetyl chloride
                                                            10400-19-8,
                         25644-88-6, S-Benzyl-L-cysteine sulfone
                                                                    82911-69-1
     Nicotinoyl chloride
     262438-43-7
                 475150-36-8
                                 701201-27-6
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
ΙT
     RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic
     preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); RACT (Reactant or reagent); USES (Uses)
        (peptide inhibitors of angiogenesis, cell migration, cell invasion and
        cell proliferation, preparation, and compns. and therapeutic uses)
RN
     701200-82-0 HCAPLUS
     L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-S-[(4-
     methylphenyl)methyl]-L-cysteinyl- (9CI) (CA INDEX NAME)
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Absolute stereochemistry.

L22 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:243058 HCAPLUS

DN 139:173332

ED Entered STN: 30 Mar 2003

TI Inhibition of integrin $\alpha 5\beta 1$ function with a small peptide (ATN-161) plus continuous 5-FU infusion reduces colorectal liver metastases and improves survival in mice

AU Stoeltzing, Oliver; Liu, Wenbiao; Reinmuth, Niels; Fan, Fan; Parry, Graham C.; Parikh, Alexander A.; McCarty, Marya F.; Bucana, Corazon D.; Mazar, Andrew P.; Ellis, Lee M.

CS Department of Cancer Biology, The University of Texas M.D. Anderson Cancer Center, Houston, TX, 77030-4009, USA

SO International Journal of Cancer (2003), 104(4), 496-503 CODEN: IJCNAW; ISSN: 0020-7136

PB Wiley-Liss, Inc.

DT Journal

LA English

AΒ

CC 1-6 (Pharmacology)

Integrin $\alpha 5\beta 1$ is expressed on activated endothelial cells and plays a critical role in tumor angiogenesis. We hypothesized that a novel integrin $\alpha 5\beta 1$ antagonist, ATN-161, would inhibit angiogenesis and growth of liver metastases in a murine model. We further hypothesized that combining ATN-161 with 5-fluorouracil (5-FU) chemotherapy would enhance the antineoplastic effect. Murine colon cancer cells (CT26) were injected into spleens of BALB/c mice to produce liver metastases. days thereafter, mice were given either ATN-161 (100 mg/kg, every 3rd day) or saline by i.p. injection, with or without combination of continuous-infusion 5-FU (100 mg/kg/2 wk), which was started on day 7. On day 20 after tumor cell inoculation, mice were killed and liver wts. and number of liver metastases were determined A follow-up study on survival was also conducted in which mice were randomized to receive ATN-161, 5-FU or ATN-161+5-FU. Combination therapy with ATN-161+5-FU significantly reduced tumor burden (liver weight) and number of liver metastases (p<0.02). Liver tumors in the ATN-161 and ATN-161+5-FU groups had significantly fewer microvessels (p<0.05) than tumors in the control or 5-FU-treated groups. Unlike treatment with either agent alone, ATN-161+5-FU significantly increased tumor cell apoptosis and decreased tumor cell proliferation (p<0.03) and improved overall survival (p<0.03, log-rank test). Targeting integrin α5β1 in combination with 5-FU infusion reduced liver metastases formation and improved survival in this colon cancer model. The enhancement of antineoplastic activity from the combination of anti-angiogenic therapy and chemotherapy may be a promising approach for treating metastatic colorectal cancer.

ST ATN 161 fluorouracil colorectal cancer integrin

IT Intestine, neoplasm

(colorectal; inhibition of integrin α5β1 function with ATN-161 plus 5-FU infusion reduces colorectal liver metastases and improves survival in mice) IT Liver, neoplasm (metastasis; inhibition of integrin $\alpha 5\beta 1$ function with ATN-161 plus 5-FU infusion reduces colorectal liver metastases and improves survival in mice) Drug interactions IT (synergistic; inhibition of integrin $\alpha 5\beta 1$ function with ATN-161 plus 5-FU infusion reduces colorectal liver metastases and improves survival in mice) IT Integrins RL: BSU (Biological study, unclassified); BIOL (Biological study) $(\alpha 5\beta 1; inhibition of integrin <math>\alpha 5\beta 1$ function with ATN-161 plus 5-FU infusion reduces colorectal liver metastases and improves survival in mice) 262438-43-7, ATN 161 IT RL: DMA (Drug mechanism of action); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (inhibition of integrin α5β1 function with ATN-161 plus 5-FU infusion reduces colorectal liver metastases and improves survival in mice) 51-21-8, 5-Fluorouracil RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL IT (Biological study); USES (Uses) (inhibition of integrin $\alpha5\beta1$ function with ATN-161 plus 5-FU infusion reduces colorectal liver metastases and improves survival in mice) THERE ARE 42 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT RE (1) Baker, C; Cancer Res 2002, V62, P1996 HCAPLUS (2) Bello, L; Cancer Res 2001, V61, P7501 HCAPLUS (3) Bergsland, E; Proc Am Soc Clin Oncol 2000, V19, P242 (4) Braakhuis, B; Semin Oncol 1995, V22, P42 HCAPLUS (5) Brooks, P; Cell 1994, V79, P1157 HCAPLUS (6) Brooks, P; J Clin Invest 1995, V96, P1815 HCAPLUS (7) Browder, T; Cancer Res 2000, V60, P1878 HCAPLUS (8) Bruns, C; Cancer 2000, V89, P488 HCAPLUS (9) Bruns, C; Clin Cancer Res 2000, V6, P1936 HCAPLUS (10) Fidler, I; Cancer Metastasis Rev 1991, V10, P229 MEDLINE (11) Friedlander, M; Science 1995, V270, P1500 HCAPLUS (12) Gately, S; Cancer J 2001, V7, P427 MEDLINE (13) Giancotti, F; Science 1999, V285, P1028 HCAPLUS (14) Gong, J; Cell Growth Differ 1997, V8, P83 HCAPLUS (15) Griggs, D; Proc Am Assoc Cancer Res 2001, V42, P1420 (16) Hanahan, D; J Clin Invest 2000, V105, P1045 HCAPLUS (17) Hynes, R; Cell 1992, V69, P11 HCAPLUS (18) Kakeji, Y; Invest New Drugs 1997, V15, P39 HCAPLUS (19) Kase, S; Anticancer Res 1993, V13, P369 HCAPLUS (20) Kerbel, R; Ann Oncol 2002, V13, P12 MEDLINE (21) Kerbel, R; Eur J Cancer 2000, V36, P1248 HCAPLUS (22) Kerr, J; Anticancer Res 1999, V19, P959 HCAPLUS (23) Kerr, J; Expert Opin Investig Drugs 2000, V9, P1271 HCAPLUS (24) Kim, S; Am J Pathol 2000, V156, P1345 HCAPLUS (25) Kim, S; J Biol Chem 2000, V275, P33920 HCAPLUS (26) Klement, G; Clin Cancer Res 2002, V8, P221 HCAPLUS (27) Klement, G; J Clin Invest 2000, V105, PR15 HCAPLUS (28) Klotz, O; Arch Clin Exp Ophthalmol 2000, V238, P88 HCAPLUS (29) Kumar, C; Adv Exp Med Biol 2000, V476, P169 MEDLINE (30) Kumar, C; Cancer Res 2001, V61, P2232 HCAPLUS (31) Livant, D; Cancer Res 2000, V60, P309 HCAPLUS (32) Lode, H; Proc Natl Acad Sci USA 1999, V96, P1591 HCAPLUS (33) Morikawa, K; J Natl Cancer Inst 1990, V82, P517 HCAPLUS (34) Mross, K; Drug Resist Updat 2000, V3, P223 HCAPLUS (35) O'Brien, V; Exp Cell Res 1996, V224, P208 HCAPLUS

(36) Remmenga, S; Gynecol Oncol 1994, V55, P115 MEDLINE

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(37) Schreiner, C; Clin Exp Metastasis 1991, V9, P163 HCAPLUS
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(38) Stoeltzing, O; Clin Cancer Res 2001, V7, P3656S

(39) Storgard, C; J Clin Invest 1999, V103, P47 HCAPLUS

(40) Tedjarati, S; Clin Cancer Res 2002, V8, P2413 HCAPLUS

(41) Varner, J; Mol Biol Cell 1995, V6, P725 HCAPLUS

(42) White, E; J Immunol 2001, V167, P5362 HCAPLUS

IT 262438-43-7, ATN 161

RL: DMA (Drug mechanism of action); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (inhibition of integrin $\alpha 5\beta 1$ function with ATN-161 plus 5-FU infusion reduces colorectal liver metastases and improves survival in mice)

RN 262438-43-7 HCAPLUS

CN L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

=> d all hitstr 124 tot

L24 ANSWER 1 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2005:409130 HCAPLUS

DN 142:458903

ED Entered STN: 13 May 2005

TI Sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy

IN Sato, Aaron K.; Sexton, Daniel J.; Dransfield, Daniel T.; Ladner, Robert C.; Arbogast, Christophe; Bussat, Philippe; Fan, Hong; Khurana, Sudha; Linder, Karen E.; Marinelli, Edmund R.; Nanjappan, Palaniappa; Nunn, Adrian D.; Pillai, Radhakrishna; Pochon, Sibylle; Ramalingam, Kondareddiar; Shrivastava, Ajay; Song, Bo; Swenson, Rolf E.; Von Wronski, Mathew A.

PA Dyax Corporation, USA; Bracco International B. V.

SO U.S. Pat. Appl. Publ., 373 pp., Cont.-in-part of U.S. Ser. No. 382,082, abandoned.

CODEN: USXXCO

DT Patent

LA English

IC ICM G01N033-53

ICS C07K007-64; A61K038-12

INCL 435007100; 514009000; 530317000

CC 6-3 (General Biochemistry)

Section cross-reference(s): 1, 3

FAN CNT 3

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	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
P.	US 2005100963	A1	20050512	US 2003-661156	20030911 <
	WO 2003074005	A2	20030912	WO 2003-US6731	20030303 <
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	GM, HR, HU,	ID, IL	, IN, IS,	JP, KE, KG, KP, KR,	KZ, LC, LK, LR,
	LS. LT. LU.	LV. MA	, MD, MG,	MK, MN, MW, MX, MZ,	NO, NZ, OM, PH,

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PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
            KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
            FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
            BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
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PRAI US 2002-360851P
    US 2003-440411P
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                                20030115
    US 2003-382082
                                20030303
                          B2
    WO 2003-US6731
                          A2
                                20030303
CLASS
PATENT NO.
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US 2005100963
                 ICM
                ICS
                        C07K007-64; A61K038-12
                 INCL
                        435007100; 514009000; 530317000
                        435/007.100; 514/009.000; 530/317.000
US 2005100963
                NCL
                        A61K049/22P4; A61K051/08Z; C07K007/08A; C07K014/00B;
                ECLA
                        C07K014/52; C07K014/71
                        A61K049/22P4; A61K051/08Z; C07K007/08A; C07K014/00B;
                ECLA
WO 2003074005
                        C07K014/52; C07K014/71
    The present invention provides polypeptides, peptide dimer, and multimeric
    complexes comprising at least one binding moiety for KDR or VEGF/KDR
    complex, which have a variety of uses wherever treating, detecting,
     isolating or localizing angiogenesis is advantageous. Particularly
    disclosed are synthetic, isolated polypeptides capable of binding KDR or
    VEGF/KDR complex with high affinity (e.g., having a KD<1 \muM), and dimer
    and multimeric constructs comprising these polypeptides.
    sequence KDR VEGF binding peptide dimer multimeric diagnosis therapy
ST
IT
    Animal cell line
        (293, 293H, as host; sequences of KDR and VEGF/KDR binding peptides,
        peptide dimers, and multimeric complexes and their use in diagnosis and
IT
     Imaging
        (NMR; sequences of KDR and VEGF/KDR binding peptides, peptide dimers,
        and multimeric complexes and their use in diagnosis and therapy)
    Animal cell line
IT
        (SW480, inhibition of; sequences of KDR and VEGF/KDR binding peptides,
        peptide dimers, and multimeric complexes and their use in diagnosis and
        therapy)
IT
    Imaging agents
        (acoustic imaging contrast agents; sequences of KDR and VEGF/KDR
        binding peptides, peptide dimers, and multimeric complexes and their
        use in diagnosis and therapy)
IT
     Imaging
        (acoustic; sequences of KDR and VEGF/KDR binding peptides, peptide
        dimers, and multimeric complexes and their use in diagnosis and
        therapy)
    Antibodies and Immunoglobulins
TT
    RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
     ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (against KDR; sequences of KDR and VEGF/KDR binding peptides, peptide
        dimers, and multimeric complexes and their use in diagnosis and
        therapy)
     Radiotherapy
IT
        (agent; sequences of KDR and VEGF/KDR binding peptides, peptide dimers,
        and multimeric complexes and their use in diagnosis and therapy)
IT
     Infection
        (bacterial; sequences of KDR and VEGF/KDR binding peptides, peptide
        dimers, and multimeric complexes and their use in diagnosis and
        therapy)
IT
     Intestine, neoplasm
        (colon, cell, inhibition of; sequences of KDR and VEGF/KDR binding
        peptides, peptide dimers, and multimeric complexes and their use in
        diagnosis and therapy)
     Escherichia coli
IT
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(enterohemorrhagic; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Gases

(fluorinated; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Protein motifs

(kinase domain region; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Proteins

RL: BSU (Biological study, unclassified); BIOL (Biological study) (linker; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Bubbles

(microbubbles, phospholipid stabilized; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Angiogenesis

(neovascularization; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Dyes

(optical; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Metals, biological studies

RL: ARG (Analytical reagent use); BSU (Biological study, unclassified); ANST (Analytical study); BIOL (Biological study); USES (Uses) (paramagnetic ion; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Molecular association

(protein-protein interaction; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Angiogenesis

Animal

Anti-AIDS agents

Antibacterial agents

Antitumor agents

Antiviral agents

Blood serum

Chelating agents

Cytotoxic agents

Drugs

Fluorescent indicators

Fluorescent substances

Human

Human immunodeficiency virus

Labels

Liposomes

Malaria

Molecular cloning

Parasiticides

Protein sequences

Radiography

Simian hemorrhagic fever virus

Simian immunodeficiency virus

cDNA library

(sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT Avidins

Perfluorocarbons

```
RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and
       multimeric complexes and their use in diagnosis and therapy)
    Enzymes, biological studies
    Radionuclides, biological studies
     RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
    ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and
       multimeric complexes and their use in diagnosis and therapy)
     Peptides, biological studies
    RL: BSU (Biological study, unclassified); DGN (Diagnostic use); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and
       multimeric complexes and their use in diagnosis and therapy)
    Imaging
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        (sonoluminescence, photoacoustic, optical; sequences of KDR and
        VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and
        their use in diagnosis and therapy)
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     Phospholipids, uses
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (stabilized microbubble; sequences of KDR and VEGF/KDR binding
       peptides, peptide dimers, and multimeric complexes and their use in
        diagnosis and therapy)
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    Particles
        (superparamagnetic; sequences of KDR and VEGF/KDR binding peptides,
        peptide dimers, and multimeric complexes and their use in diagnosis and
       therapy)
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    AIDS (disease)
     Neoplasm
        (treatment of; sequences of KDR and VEGF/KDR binding peptides, peptide
        dimers, and multimeric complexes and their use in diagnosis and
        therapy)
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     Vascular endothelial growth factor receptors
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (type VEGFR-2; sequences of KDR and VEGF/KDR binding peptides, peptide
        dimers, and multimeric complexes and their use in diagnosis and
        therapy)
IT
     Infection
        (viral; sequences of KDR and VEGF/KDR binding peptides, peptide dimers,
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        (amino acid sequence; sequences of KDR and VEGF/KDR binding peptides,
        peptide dimers, and multimeric complexes and their use in diagnosis and
        therapy)
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     (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (amino acid sequence; sequences of KDR and VEGF/KDR binding peptides,
        peptide dimers, and multimeric complexes and their use in diagnosis and
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     76-16-4, Perfluoroethane 76-19-7, Perfluoropropane
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     Bromochlorodifluoromethane 355-25-9, Perfluorobutane
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203Pb, uses 14701-22-5, Ni2+, uses 14733-03-0, 214Bi, uses
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     15743-54-1, 168Yb, uses 15750-15-9, 111In, uses
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69146-59-4, MECAM 71353-07-6, LICAM 79642-50-5 104162-48-3, DOTMA 114873-37-9, DO3A 157885-16-0, NeutrAvidin 325126-08-7, TETMA RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses) (sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT 127464-60-2, VEGF

ΙT

RL: BSU (Biological study, unclassified); BIOL (Biological study) (sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

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RL: PRP (Properties)

(unclaimed nucleotide sequence; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT 851737-28-5 851737-29-6

RL: PRP (Properties)

(unclaimed protein sequence; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

IT 599210-01-2 851724-81-7

RL: BSU (Biological study, unclassified); DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (amino acid sequence; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

RN 599210-01-2 HCAPLUS

CN L-Glutamic acid, glycyl-L-histidyl-L-prolyl-L-cysteinyl-L-lysylglycyl-L-methionyl-L-leucyl-L-prolyl-L-histidyl-L-threonyl-L-cysteinyl-L-tryptophyl-L-tyrosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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PAGE 1-B

RN 851724-81-7 HCAPLUS
CN L-Lysine, L-alanylglycylglycyl-L-histidyl-L-prolyl-L-cysteinyl-L-lysylglycyl-L-methionyl-L-leucyl-L-prolyl-L-histidyl-L-threonyl-L-cysteinyl-L-tryptophyl-L-tyrosyl-L-α-glutamylglycyl-L-threonylglycylglycylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

PAGE 1-C

PAGE 2-C

L24 ANSWER 2 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2005:238537 HCAPLUS

DN 142:310938

ED Entered STN: 18 Mar 2005

Fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes (phenol oxidase B or laccase), and their use in construction of detergent for bleaching carotenoid stain on fabric or

Search done by Noble Jarrell

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surface
     Wolfgang, Aehle; Baldwin, Toby M.; Van Gastel, Franciscus J. C.; Janssen,
IN
     Giselle G.; Murray, Christopher J.; Wang, Huaming; Winetzky, Deborah S.
PΑ
     U.S. Pat. Appl. Publ., 107 pp.
SO
     CODEN: USXXCO
DТ
     Patent
LÀ
     English
     ICM C12Q001-68
IC
     ICS C07H021-04; C12N009-02; C11D003-386
INCL 435006000; 435069100; 435189000; 435320100; 435325000; 536023200;
     510320000
     3-2 (Biochemical Genetics)
     Section cross-reference(s): 7, 10, 46
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                                            APPLICATION NO.
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                                20050317
                                          US 2002-235043
     US 2005058996
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PΙ
                                20020903 <--
PRAI US 2002-235043
CLASS
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 US 2005058996
                 ICM
                        C07H021-04; C12N009-02; C11D003-386
                 ICS
                        435006000; 435069100; 435189000; 435320100; 435325000;
                 INCL
                        536023200; 510320000
                        435/006.000; 435/069.100; 435/189.000; 435/320.100;
 US 2005058996
                 NCL
                        435/325.000; 536/023.200; 510/320.000
                        C11D003/386H; C12N009/02H3B2
                 ECLA
     The invention provides numerous peptides composed of 7 or 12 amino acids
AB
     that are able to bind to a carotenoid compound on a fabric. The invention
     relates said peptides: (a) may also contain a cysteine added to each end
     (defined as C-C derivs.); (b) comprise a repeatable motif; and (c) bind to
     tomato or paprika stains on cotton. The invention also provides chimeric
     proteins composed of said peptides linked to the C-terminus of a phenol
     oxidizing enzyme, such as laccase and/or Stachybotrys phenol oxidase B,
     and/or variants thereof. The invention further provides polynucleotides
     encoding said carotenoid-binding peptide-phenol oxidizing enzyme fusion
     proteins, and use of said polynucleotides in transforming host cells for recombinant production of fusion proteins. Still further, the invention
     provides for the use of said recombinant fusion proteins in construction
     of a detergent, wherein said detergent can be used to clean a carotenoid
     stain on a fabric and/or a surface contacting the stain. Finally, the
     invention provides the amino acid sequences of Stachybotrys chartarum
     phenol oxidase B and carotenoid-binding peptides. In the examples, the
     invention demonstrated that a fusion protein composed of Seq ID number 16
     carotenoid-binding peptide and a phenol oxidase B variant
     (M254F/E346V/E348Q) bound to tomato stain on cotton better than the enzyme
     alone.
     fusion protein carotenoid binding peptide phenol oxidizing enzyme;
ST
     sequence phenol oxidase B Stachybotrys use fusion protein; detergent stain
     remover carotenoid binding peptide phenol oxidizing enzyme; laccase
     carotenoid binding peptide fusion detergent stain remover
IT
     Detergents
        (bleaching; fusion proteins composed of carotenoid-binding peptides and
        phenol oxidizing enzymes, and their use in construction of detergent
        for bleaching carotenoid stain on fabric or surface)
IT
     Fusion proteins (chimeric proteins)
     RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or
     engineered material use); BIOL (Biological study); PREP (Preparation);
     USES (Uses)
        (carotenoid-binding peptide fused to phenol oxidizing enzyme; fusion
        proteins composed of carotenoid-binding peptides and phenol oxidizing
        enzymes, their polynucleotides and use in construction of detergent)
IT
     Peptides, properties
     RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or
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engineered material use); BIOL (Biological study); PREP (Preparation); USES (Uses) (carotenoid-binding, fusion product with phenol oxidizing enzyme; fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, and their use in construction of detergent) Textiles (cotton; fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, and their ability to better bind tomato stains on cotton than enzymes alone) Detergents Surface Textiles (fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, and their use in construction of detergent for bleaching carotenoid stain on fabric or surface) Carotenes, processes RL: REM (Removal or disposal); PROC (Process) (fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, and their use in construction of detergent for bleaching carotenoid stain on fabric or surface) Polynucleotides RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses) (fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, their polynucleotides and use in construction of detergent for bleaching carotenoid stain on fabric or surface) Capsicum annuum annuum (longum group, paprika; peptides able to bind carotenoid compds., their ability to bind carotenoids found in tomato or paprika and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes) Plasmid vectors (pGAPT; polynucleotides encoding fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, their vectors and use in transforming host cells for recombinant protein production) Lycopersicon esculentum (peptides able to bind carotenoid compds., their ability to bind carotenoids found in tomato or paprika and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes) Repeat motifs (protein) (peptides able to bind carotenoid compds., their protein motifs, sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes for creation of detergent) Molecular cloning (polynucleotides encoding fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, their vectors and use in transforming host cells for recombinant protein production) Stachybotrys chartarum (sequence of phenol oxidase B found in Stachybotrys chartarum, and its use in production of fusion proteins comprising carotenoid-binding proteins) Detergents (stain removers; fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, and their use in construction of detergent for bleaching carotenoid stain on fabric or surface) Aspergillus niger

Escherichia coli

(transformed; polynucleotides encoding fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, their vectors and use in transforming host cells for recombinant protein production)

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RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or engineered material use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(amino acid sequence; fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes, and their ability to better bind ΙT

tomato stains on cotton than enzymes alone) 502164-47-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-48-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-50-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-52-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-53-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-54-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-55-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-56-9DP, cysteine derivs. and fusion product with phenol oxidizing 502164-58-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-59-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-60-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-61-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-62-7DP, cysteine 502164-63-8DP, derivs. and fusion product with phenol oxidizing enzyme cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-64-9DP, cysteine derivs. and fusion product with phenol oxidizing 502164-65-0DP, cysteine derivs. and fusion product with phenol enzyme oxidizing enzyme 502164-66-1DP, cysteine derivs. and fusion product with 502164-67-2DP, cysteine derivs. and fusion phenol oxidizing enzyme 502164-68-3DP, cysteine derivs. and product with phenol oxidizing enzyme fusion product with phenol oxidizing enzyme 502164-69-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-70-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-71-8DP, cysteine derivs. and fusion product with phenol oxidizing 502164-72-9DP, cysteine derivs. and fusion product with phenol enzvme oxidizing enzyme 502164-73-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-74-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-75-2DP, cysteine derivs. and 502164-76-3DP, cysteine fusion product with phenol oxidizing enzyme derivs. and fusion product with phenol oxidizing enzyme 502164-77-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-78-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-79-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-80-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-81-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-83-2DP, cysteine derivs. and 502164-84-3DP, cysteine fusion product with phenol oxidizing enzyme derivs. and fusion product with phenol oxidizing enzyme 502164-85-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-86-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-87-6DP, cysteine derivs. and fusion product with phenol 502164-88-7DP, cysteine derivs. and fusion product with oxidizing enzyme 502164-89-8DP, cysteine derivs. and fusion phenol oxidizing enzyme product with phenol oxidizing enzyme 502164-90-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-91-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-92-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-93-4DP, cysteine derivs. and fusion product with phenol oxidizing 502164-94-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-95-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-96-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-97-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-98-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502164-99-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-00-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-02-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-04-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-06-2DP, cysteine derivs. and fusion 502165-08-4DP, cysteine derivs. and product with phenol oxidizing enzyme 502165-10-8DP, cysteine fusion product with phenol oxidizing enzyme derivs. and fusion product with phenol oxidizing enzyme 502165-11-9DP. cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-13-1DP, cysteine derivs. and fusion product with phenol oxidizing

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502165-80-2DP, cysteine derivs. and fusion phenol oxidizing enzyme product with phenol oxidizing enzyme 502165-81-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-82-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-83-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-84-6DP, cysteine derivs. and fusion product with phenol oxidizing 502165-85-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-86-8DP, cysteine derivs. and fusion product with 502165-87-9DP, cysteine derivs. and fusion phenol oxidizing enzyme 502165-88-0DP, cysteine derivs. and product with phenol oxidizing enzyme fusion product with phenol oxidizing enzyme 502165-89-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-90-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-91-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-92-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-93-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-94-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-95-9DP, cysteine derivs. and 502165-96-0DP, cysteine fusion product with phenol oxidizing enzyme derivs. and fusion product with phenol oxidizing enzyme 502165-97-1DP. cysteine derivs. and fusion product with phenol oxidizing enzyme 502165-98-2DP, cysteine derivs. and fusion product with phenol oxidizing 502165-99-3DP, cysteine derivs. and fusion product with phenol enzvme oxidizing enzyme 502166-00-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-01-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-02-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-03-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-04-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-05-4DP, cysteine derivs. and fusion product with phenol oxidizing 502166-06-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-07-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-09-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-10-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-11-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-12-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-13-4DP, cysteine derivs. and fusion product with phenol oxidizing 502166-14-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-15-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-16-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-17-8DP, cysteine derivs. and 502166-18-9DP, cysteine fusion product with phenol oxidizing enzyme derivs. and fusion product with phenol oxidizing enzyme 502166-19-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-20-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-21-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-22-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-23-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-24-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-25-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-26-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-27-0DP, cysteine derivs. and fusion product with phenol oxidizing 502166-28-1DP, cysteine derivs. and fusion product with phenol 502166-29-2DP, cysteine derivs. and fusion product with oxidizing enzyme phenol oxidizing enzyme 502166-30-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-31-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-32-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-34-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-35-0DP, cysteine derivs. and fusion product with phenol oxidizing 502166-36-1DP, cysteine derivs. and fusion product with phenol enzyme oxidizing enzyme 502166-37-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-38-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-39-4DP, cysteine derivs. and

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enzyme 502166-97-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-98-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502166-99-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-00-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or engineered material use); BIOL (Biological study); PREP (Preparation); USES (Uses)

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(amino acid sequence; peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes for creation of detergent) 502167-01-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-02-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-03-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-04-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-05-7DP, cysteine derivs. and 502167-06-8DP, cysteine fusion product with phenol oxidizing enzyme 502167-07-9DP, derivs. and fusion product with phenol oxidizing enzyme cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-08-0DP, cysteine derivs. and fusion product with phenol oxidizing 502167-09-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-10-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-11-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-12-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-13-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-14-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-15-9DP, cysteine derivs. and fusion product with phenol oxidizing 502167-16-0DP, cysteine derivs. and fusion product with phenol enzvme oxidizing enzyme 502167-17-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-18-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-19-3DP, cysteine derivs. and 502167-20-6DP, cysteine fusion product with phenol oxidizing enzyme derivs. and fusion product with phenol oxidizing enzyme 502167-21-7DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-22-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-23-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-24-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-25-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-26-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-27-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-29-5DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-30-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-31-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-32-0DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-33-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-34-2DP, cysteine derivs. and 502167-35-3DP, cysteine fusion product with phenol oxidizing enzyme derivs. and fusion product with phenol oxidizing enzyme 502167-36-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-37-5DP, cysteine derivs. and fusion product with phenol oxidizing 502167-38-6DP, cysteine derivs. and fusion product with phenol 502167-39-7DP, cysteine derivs. and fusion product with oxidizing enzyme 502167-40-0DP, cysteine derivs. and fusion phenol oxidizing enzyme product with phenol oxidizing enzyme 502167-41-1DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-42-2DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-43-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-44-4DP, cysteine derivs. and fusion product with phenol oxidizing 502167-45-5DP, cysteine derivs. and fusion product with phenol enzyme oxidizing enzyme 502167-46-6DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-47-7DP, cysteine derivs. and fusion

product with phenol oxidizing enzyme 502167-48-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-49-9DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-50-2DP,

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(amino acid sequence; peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes for creation of detergent)

847966-55-6DP, fusion product with carotenoid-binding peptide RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or engineered material use); BIOL (Biological study); PREP (Preparation); USES (Uses)

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IT

(amino acid sequence; sequence of phenol oxidase B found in Stachybotrys chartarum, and its use in production of fusion proteins comprising carotenoid-binding proteins)

9002-10-2DP, Phenol oxidase, fusion product with carotenoid-binding peptide

RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or engineered material use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(isoenzyme B; peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes (such as phenol oxidase B) for creation of detergent) 80498-15-3DP, Laccase, fusion product with carotenoid-binding peptide RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or engineered material use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing

enzymes (such as laccase) for creation of detergent)
IT 52-90-4DP, Cysteine, added to N-terminus or C-terminus of carotenoid-binding peptide

RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or engineered material use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes for creation of detergent)

IT 502697-64-5 502697-65-6 847967-48-0 847967-49-1 847967-50-4 847967-51-5 847967-52-6 847967-53-7 847967-54-8 847967-55-9 847967-56-0 847967-57-1 847967-58-2 RL: PRP (Properties)

(unclaimed sequence; fusion proteins composed of carotenoid-binding peptides and phenol oxidizing enzymes (phenol oxidase B or laccase), and their use in construction of detergent for bleaching carotenoid stain on fabric or surface)

502166-83-8DP, cysteine derivs. and fusion product with phenol
 oxidizing enzyme 502167-78-4DP, cysteine derivs. and fusion
 product with phenol oxidizing enzyme 502167-93-3DP, cysteine
 derivs. and fusion product with phenol oxidizing enzyme
 RL: BPN (Biosynthetic preparation); PRP (Properties); TEM (Technical or
 engineered material use); BIOL (Biological study); PREP (Preparation);
 USES (Uses)

(amino acid sequence; peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes for creation of detergent)

RN 502166-83-8 HCAPLUS

CN L-Methionine, L-isoleucyl-L-asparaginyl-L-threonyl-L-prolyl-L-histidyl-L-seryl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 502167-78-4 HCAPLUS

CN L-Proline, L-seryl-L-prolyl-L-histidyl-L-seryl-L-methionyl-L-leucyl-L-glutaminyl-L-asparaginyl-L-prolyl-L-serylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

RN 502167-93-3 HCAPLUS

CN L-Tryptophan, L-valyl-L-alanyl-L-seryl-L-alanyl-L-asparaginyl-L-prolyl-L-histidyl-L-seryl-L-methionyl-L-threonyl-L-seryl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B

PAGE 2-B

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ANSWER 3 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
     2004:634089 HCAPLUS
AN
      141:167846
ED
     Entered STN: 06 Aug 2004
     Binding peptides for the KDR receptor and vascular endothelial growth
      factor/KDR complex and their use in diagnosis, therapy, and imaging of
      angiogenesis-related disorders
IN
     Sato, Aaron K.; Sexton, Daniel J.; Dransfield, Daniel T.; Ladner, Robert
     C.; Arbogast, Christophe; Bussat, Philippe; Fan, Hong; Khurana, Sudha;
     Linder, Karen E.; Marinelli, Edmund R.; Nanjappan, Palaniappa; Nunn,
     Adrian; Pillai, Radhakrishna; Pochon, Sibylle; Ramalingam, Kondareddiar; Shrivastava, Ajay; Song, Bo; Swenson, Rolf E.; Von Wronski, Mathew A.
     Dyax Corp., USA; Bracco International B.V.
PA
     PCT Int. Appl., 470 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LΑ
      English
      ICM C12Q001-00
IC
     ICS A61K048-00
      1-12 (Pharmacology)
      Section cross-reference(s): 9, 63
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                                                 APPLICATION NO.
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      PATENT NO.
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PΙ
     WO 2004065621
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                                    20040805
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                                 20030115
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CLASS
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                         A61K048-00
                  ICS
                         A61K049/22P4; A61K051/08Z; C07K007/08A; C07K014/00B;
 WO 2004065621
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                         C07K014/52; C07K014/71
WO 2003074005
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                         A61K049/22P4; A61K051/08Z; C07K007/08A; C07K014/00B;
                         C07K014/52; C07K014/71
     MARPAT 141:167846
os
     The present invention provides peptides, peptide dimers, and multimeric
AΒ
     complexes comprising at least one binding moiety for KDR receptor or
     vascular endothelial growth factor (VEGF)/KDR complex, which have a
     variety of uses wherever treating, detecting, isolating, or localizing
     angiogenesis is advantageous. Particularly disclosed are synthetic, isolated peptides capable of binding KDR or VEFG/KDR complex with high
     affinity (e.g., having a KD < 1 \mu M) , and dimer and multimeric
     constructs comprising these polypeptides. The involvement of VEGF and KDR
     in angiogenesis makes the binding peptides particularly useful for imaging
     important sites of angiogenesis, e.g., neoplastic tumors, for targeting
     substances, e.g., therapeutics, including radiotherapeutics, to such
     sites, and for treating certain disease states, including those associated
     with inappropriate angiogenesis.
     KDR receptor VEGF binding peptide angiogenesis diagnosis therapy; vascular
     endothelial growth factor KDR receptor binding peptide; imaging KDR
     receptor VEGF binding peptide angiogenesis
IT
     Imaging agents
        (NMR contrast; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
     Imaging agents
        (acoustic imaging contrast agents; binding peptides for the KDR
        receptor and vascular endothelial growth factor/KDR complex and their
        use in diagnosis, therapy, and imaging of angiogenesis-related
IT
     Peptides, biological studies
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (amides; binding peptides for the KDR receptor and vascular endothelial
        growth factor/KDR complex and their use in diagnosis, therapy, and
        imaging of angiogenesis-related disorders)
     Peptides, biological studies
ΙT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (analogs, peptoids; binding peptides for the KDR receptor and vascular
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endothelial growth factor/KDR complex and their use in diagnosis,

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therapy, and imaging of angiogenesis-related disorders)
IT
     Angiogenesis inhibitors
     Anti-AIDS agents
     Antimalarials
     Antitumor agents
     Antiviral agents
     Chemotherapy
     Drug screening
     Human
     Neoplasm
     Phage display
     Radiopharmaceuticals
     Radiotherapy
     Scintigraphic agents
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
IT
     Avidins
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
     Glycopeptides
TT
     Peptides, biological studies
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
IT
     Chelating agents
     Cytotoxic agents
     Dves
     Fluorescent substances
     Paramagnetic materials
        (conjugates with peptides; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
     Radionuclides, biological studies
TT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (conjugates with peptides; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
     Enzymes, biological studies
TT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (conjugates, with peptides; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
     Peptides, biological studies
ΙT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
         (conjugates; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
         (contrast, radiog.; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
     Human immunodeficiency virus
TT
     Malaria
     Simian hemorrhagic fever virus
     Simian immunodeficiency virus
         (detection of; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
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therapy, and imaging of angiogenesis-related disorders)
IT
     Imaging
        (diagnostic; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
     Peptides, biological studies
TΤ
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (disulfide-containing; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
тт
     Escherichia coli
        (enterohemorrhagic, detection of; binding peptides for the KDR receptor
        and vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
IT
        (fluoronated, conjugates with peptides; binding peptides for the KDR
        receptor and vascular endothelial growth factor/KDR complex and their
        use in diagnosis, therapy, and imaging of angiogenesis-related
        disorders)
     Angiogenesis
IT
        (inhibition of; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
     Peptides, biological studies
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (labeled; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
     Drug delivery systems
        (liposomes, peptide-containing; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
IT
     Bubbles
        (microbubbles, phospholipid-stabilized, conjugates with peptides;
        binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
IT
     Diagnosis
        (mol.; binding peptides for the KDR receptor and vascular endothelial
        growth factor/KDR complex and their use in diagnosis, therapy, and
        imaging of angiogenesis-related disorders)
IT
        (paramagnetic, conjugates with peptides; binding peptides for the KDR
        receptor and vascular endothelial growth factor/KDR complex and their
        use in diagnosis, therapy, and imaging of angiogenesis-related
        disorders)
TT
     Bacteriophage
        (peptide display; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
        (peptide-containing; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
     Peptides, biological studies
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (retro-inverso; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
     Paramagnetic materials
        (superparamagnetic, conjugates with peptides; binding peptides for the
        KDR receptor and vascular endothelial growth factor/KDR complex and
        their use in diagnosis, therapy, and imaging of angiogenesis-related
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disorders)
IT
    Drug delivery systems
    Gene targeting
        (to KDR-expressing cells; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
    Vascular endothelial growth factor receptors
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
    use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological
     study); USES (Uses)
        (type VEGFR-2; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
     Perfluorocarbons
     RL: DGN (Diagnostic use); BIOL (Biological study); USES (Uses)
        (ultrasound contrast agent; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
     612494-38-9
IT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial 1growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
     127464-60-2D, Vascular endothelial growth factor, KDR receptor complexes
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological
     study); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
     58-85-5, Biotin 9013-20-1, Streptavidin
                                                 157885-16-0, Neutravidin
IT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
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     736140-24-2P
                    736140-25-3P
TT
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                    736140-32-2P
                                   736140-33-3P
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                    736141-90-5P
                                   736141-92-7P
     736141-87-0P
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); SPN (Synthetic
     preparation); THU (Therapeutic use); ANST (Analytical study); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
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                                 599208-47-6
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RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
   (binding peptides for the KDR receptor and vascular endothelial growth
   factor/KDR complex and their use in diagnosis, therapy, and imaging of
   angiogenesis-related disorders)
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                                                           599211-74-2
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733170-07-5D, conjugates with biotin
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    RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
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     736140-23-1
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (binding peptides for the KDR receptor and vascular endothelial growth
        factor/KDR complex and their use in diagnosis, therapy, and imaging of
        angiogenesis-related disorders)
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TT
     189023-89-0 444812-91-3 612493-70-6
     RL: DGN (Diagnostic use); BIOL (Biological study); USES (Uses)
        (chelator; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
     51-92-3, Tetramethylammonium 60-00-4, EDTA, biological studies
TТ
                    112-24-3, TETA 869-52-3, TTHA 1170-02-1
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     67-43-6, DTPA
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                                     104162-48-3, DOTMA
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     RL: DGN (Diagnostic use); THU (Therapeutic use); BIOL (Biological study);
     USES (Uses)
        (chelator; binding peptides for the KDR receptor and vascular
        endothelial growth factor/KDR complex and their use in diagnosis,
        therapy, and imaging of angiogenesis-related disorders)
IT
     735845-12-2
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (consensus peptide sequence 10; binding peptides for the KDR receptor
        and vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
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TT 735845-13-3
RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)

IT

IT

- use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (consensus peptide sequence 11; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders) 735845-14-4
- RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (consensus peptide sequence 12; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders) 735845-15-5
- RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (consensus peptide sequence 14; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders) 735845-07-5
- IT 735845-07-5
 RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
 (consensus peptide sequence 5; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders)
 IT 735845-10-0
 - RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (consensus peptide sequence 7; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders) 735845-11-1
- TT 735845-11-1
 RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
 (consensus peptide sequence 9; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders)
- TT 735845-16-6
 RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
 (loop consensus peptide sequence 13; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders)
- TT 735845-18-8
 RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
 (loop consensus peptide sequence 24; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders)
- IT 735845-19-9
 RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
 (loop consensus peptide sequence 25; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders)
- IT 14701-22-5, Nickel-2+, biological studies 14913-52-1, Neodymium-3+, biological studies 15158-11-9, Copper-2+, biological studies

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16065-83-1, Chromium-3+,
    15438-31-0, Iron-2+, biological studies
                        16397-91-4, Manganese-2+, biological studies
    biological studies
    16910-54-6, Europium-2+, biological studies 18472-30-5, Erbium-3+,
    biological studies 20074-52-6, Iron-3+, biological studies 22537-59-3,
    Protactinium-4+, biological studies 22541-14-6, Praseodymium-3+,
    biological studies 22541-17-9, Samarium-3+, biological studies
     22541-18-0, Europium-3+, biological studies
                                                   22541-19-1, Gadolinium-3+,
                         22541-20-4, Terbium-3+, biological studies
    biological studies
    22541-21-5, Dysprosium-3+, biological studies 22541-22-6, Holmium-3+,
    biological studies
                         22541-53-3, Cobalt-2+, biological studies
                                               22541-75-9, Titanium-3+,
     22541-63-5, Cobalt-3+, biological studies
    biological studies
     RL: DGN (Diagnostic use); THU (Therapeutic use); BIOL (Biological study);
    USES (Uses)
        (paramegnaetic metal; binding peptides for the KDR receptor and
       vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
     10043-49-9, Gold-198, biological studies 10043-66-0, Iodine-131,
TT
                         10098-91-6, Yttrium-90, biological studies
     biological studies
     13967-64-1, Dysprosium-165, biological studies
                                                     13967-65-2, Holmium-166,
                         13967-74-3, Cerium-141, biological studies
     biological studies
    13968-53-1, Ruthenium-103, biological studies 13981-25-4, Copper-64, biological studies 13981-28-7, Lanthanum-140, biological studies
     13981-56-1, Fluorine-18, biological studies 13982-36-0, Yttrium-88,
                        14041-44-2, Ytterbium-175, biological studies
     biological studies
     14119-09-6, Gallium-67, biological studies 14158-30-6, Iodine-124,
                        14158-31-7, Iodine-125, biological studies
     biological studies
     14265-75-9, Lutetium-177, biological studies
                                                   14276-53-0, Copper-62,
                        14378-26-8, Rhenium-188, biological studies
     biological studies
                                              14391-19-6, Terbium-161,
     14391-11-8, Gold-199, biological studies
                        14391-22-1, Thulium-167, biological studies
     biological studies
     14391-96-9, Scandium-47, biological studies
                                                   14392-02-0, Chromium-51,
                         14687-25-3, Lead-203, biological studies
     biological studies
     14733-03-0, Bismuth-214, biological studies
                                                  14913-49-6, Bismuth-212,
     biological studies 14913-89-4, biological studies 14981-64-7,
     Palladium-109, biological studies 14998-63-1, Rhenium-186, biological
             15229-37-5, Bismuth-211, biological studies 15715-08-9,
     Iodine-123, biological studies
                                    15743-54-1, Ytterbium-168, biological
              15750-15-9, Indium-111, biological studies 15757-14-9,
     Gallium-68, biological studies 15757-86-5, Copper-67, biological studies
     15758-35-7, Ruthenium-97, biological studies 15765-31-8, Promethium-149,
     biological studies 15765-38-5, Bromine-76, biological studies
     15765-39-6, Bromine-77, biological studies 15766-00-4, Samarium-153,
                         15776-20-2, Bismuth-213, biological studies
     biological studies
     15840-01-4, Dysprosium-166, biological studies
                                                     378784-45-3,
     Technetium-99m, biological studies
     RL: DGN (Diagnostic use); THU (Therapeutic use); BIOL (Biological study);
     USES (Uses)
        (therapeutic radionuclide; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
     75-71-8, Dichlorodifluoromethane
                                        75-72-9, Chlorotrifluoromethane
ΤT
                                  76-15-3
                                            76-19-7, Octafluoropropane
     75-73-0, Tetrafluoromethane
     353-59-3, Bromochlorodifluoromethane
                                            355-25-9, Decafluorobutane
     RL: DGN (Diagnostic use); BIOL (Biological study); USES (Uses)
        (ultrasound contrast agent; binding peptides for the KDR receptor and
        vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
                                 735863-60-2
                                               735863-61-3
                                                             735863-62-4
                   735863-59-9
     735863-58-8
ΙT
                                 735863-65-7
                                               735863-66-8
     735863-63-5
                   735863-64-6
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; binding peptides for the KDR receptor
        and vascular endothelial growth factor/KDR complex and their use in
        diagnosis, therapy, and imaging of angiogenesis-related disorders)
                                 599210-03-4
                                               599210-04-5
                                                             735281-23-9
     210573-59-4
                   599210-02-3
IT
                                               735863-68-0
                                 735863-67-9
                   735281-25-1
     735281-24-0
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RL: PRP (Properties)

(unclaimed sequence; binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders)

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD

- RE (1) Hirth; US 5942385 A 1999 HCAPLUS
- (2) Hu; US 6040157 A 2000 HCAPLUS
- (3) Keyt; US 6057428 A 2000 HCAPLUS
- (4) Ullrich; US 5851999 A 1998 HCAPLUS
- IT 599210-01-2

CN

RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (binding peptides for the KDR receptor and vascular endothelial growth factor/KDR complex and their use in diagnosis, therapy, and imaging of angiogenesis-related disorders)

RN 599210-01-2 HCAPLUS

L-Glutamic acid, glycyl-L-histidyl-L-prolyl-L-cysteinyl-L-lysylglycyl-L-methionyl-L-leucyl-L-prolyl-L-histidyl-L-threonyl-L-cysteinyl-L-tryptophyl-L-tyrosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

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L24 ANSWER 4 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
     2004:311016 HCAPLUS
AN
DN
     140:337904
     Entered STN: 16 Apr 2004
ED
     Vaccines comprising optimized multiple cytotoxic or helper T lymphocyte
ΤI
     epitopes derived from hepatitis B virus pol, env and core proteins
     Sette, Alessandro; Chesnut, Robert; Newman, Mark J.; Livingston, Brian D.;
IN
     Babe, Lilia Maria; Chen, Yiyou; Deyoung, Lawrence M.; Huang, Manley T. F.;
     Power, Scott D.
     Epimmune Inc., USA; Genencor International, Inc.
PA
     PCT Int. Appl., 401 pp.
so
      CODEN: PIXXD2
DT
     Patent
     English
LА
IC
     ICM C07K
     15-2 (Immunochemistry)
CC
     Section cross-reference(s): 3
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     PATENT NO.
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WO 2004031210

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Search done by Noble Jarrell

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,

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CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
              LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ,
              OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
              TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
              BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     CA 2500955
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PRAI US 2002-415463P
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     US 2002-419973P
                            ₽
                                  20021022
                                             <--
     WO 2003-US31303
                            W
                                  20031003
CLASS
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                         C07K
 WO 2004031210
                  ICM
     The invention relates to multi-epitope nucleic acid and peptide vaccines
AΒ
     and methods of designing such vaccines to provide increased
     immunogenicity. In particular, the chimeric constructs comprise cytotoxic
     T lymphocyte and helper T cell epitopes derived from hepatitis B virus
     pol, env and core proteins.
     hepatitis B virus pol env core protein CTL epitope; cytotoxic T lymphocyte
ST
     epitope HBV antigen hepatitis B vaccine; helper T cell epitope HBcAg pol
     env HBV vaccine
     Histocompatibility antigens
TТ
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (HLA (human leukocyte-associated antigen); vaccines comprising optimized
        multiple cytotoxic or helper T lymphocyte epitopes derived from
        hepatitis B virus pol, env and core proteins)
TΤ
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (HLA-A1; vaccines comprising optimized multiple cytotoxic or helper T
         lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
TТ
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (HLA-A24; vaccines comprising optimized multiple cytotoxic or helper T
         lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
IT
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (HLA-A2; vaccines comprising optimized multiple cytotoxic or helper T
         lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
     Histocompatibility antigens
IT
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (HLA-A3; vaccines comprising optimized multiple cytotoxic or helper T
         lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
TT
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
         (HLA-B, HLA-B3; vaccines comprising optimized multiple cytotoxic or
         helper T lymphocyte epitopes derived from hepatitis B virus pol, env
         and core proteins)
     Histocompatibility antigens
IT
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
         (HLA-D, HLA-DM; vaccines comprising optimized multiple cytotoxic or
         helper T lymphocyte epitopes derived from hepatitis B virus pol, env
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and core proteins)
IT
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (HLA-D, HLA-DO; vaccines comprising optimized multiple cytotoxic or
        helper T lymphocyte epitopes derived from hepatitis B virus pol, env
        and core proteins)
TΤ
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (HLA-DR3; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
     Histocompatibility antigens
IT
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (HLA-DR; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
IT
     Genetic element
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (IRES (internal ribosomal entry site) element; vaccines comprising
        optimized multiple cytotoxic or helper T lymphocyte epitopes derived
        from hepatitis B virus pol, env and core proteins)
IT
     Proteins
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (LAMP-1 (lysosome-associated membrane protein 1), lysosomal targeting
        sequence; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
IT
     Proteins
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (LAMP-2 (lysosome-associated membrane protein 2), lysosomal targeting
        sequence; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
     Histocompatibility antigens
TT .
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (MHC (major histocompatibility complex), class I, targeting; vaccines
        comprising optimized multiple cytotoxic or helper T lymphocyte epitopes
        derived from hepatitis B virus pol, env and core proteins)
     Histocompatibility antigens
TТ
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (MHC (major histocompatibility complex), class II, targeting; vaccines
        comprising optimized multiple cytotoxic or helper T lymphocyte epitopes
        derived from hepatitis B virus pol, env and core proteins)
TΤ
     Proteins
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (Ty; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
IT
     Prostate gland, neoplasm
        (antigen; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
IT
        (cell; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
     Protein motifs
```

IT

(cytoplasmic domain; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT T cell (lymphocyte)

(cytotoxic; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT T cell (lymphocyte)

(helper cell; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Antigens

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(hepatitis B core; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Antigens

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(hepatitis B surface; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Vaccines

(hepatitis B; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Vaccines

(influenza; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Influenza

(matrix protein; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Proteins

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(matrix, influenza; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Polyproteins

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(pol; vaccines comprising optimized multiple cytotoxic or helper T .lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Cytomegalovirus

(promoter; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Genetic element

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(regulatory; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env and core proteins)

IT Genetic element

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(signal sequence; vaccines comprising optimized multiple cytotoxic or helper T lymphocyte epitopes derived from hepatitis B virus pol, env

and core proteins)
IT Lysosome

(targeting sequence; vaccines comprising optimized multiple cytotoxic

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or helper T lymphocyte epitopes derived from hepatitis B virus pol, env
        and core proteins)
IT
    DNA
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (vaccine; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
IT
     DNA sequences
     Endoplasmic reticulum
     Epitopes
     Gene therapy
     Genetic vectors
     Hepatitis B virus
     Hepatitis C virus
     Human
     Human immunodeficiency virus
     Immunotherapy
     Molecular cloning
     Protein sequences
        (vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
IT
     Envelope proteins
     Fusion proteins (chimeric proteins)
     Nucleic acids
     Polynucleotides
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
     Invariant chain (class II antigen)
     Promoter (genetic element)
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
IT
     Antibodies and Immunoglobulins
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (κ; vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
                                                   679448-52-3P
                                                                  679448-53-4P
IT
     679448-45-4P
                    679448-46-5P
                                   679448-51-2P
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                    679449-09-3P
                                   679449-10-6P
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     679449-08-2P
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                                                                  679449-24-2P
     679449-20-8P
                    679449-21-9P
                                   679449-22-0P
                    679449-26-4P
     679449-25-3P
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (amino acid sequence; vaccines comprising optimized multiple cytotoxic
        or helper T lymphocyte epitopes derived from hepatitis B virus pol, env
        and core proteins)
                                                   679448-48-7P
                                                                  679448-49-8P
ΙT
     679448-43-2P
                    679448-44-3P
                                   679448-47-6P
                    679448-55-6P
     679448-50-1P
                                   679448-56-7P
                                                   679448-57-8P
                                                                  679448-58-9P
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679448-62-5P
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     679448-59-0P
                    679448-60-3P
                                    679448-61-4P
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                                                                   679448-68-1P
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                                                                   679448-78-3P
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                    679448-80-7P
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                                                   679449-16-2P
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     679449-13-9P
                    679449-14-0P
     679449-18-4P
                  . 679449-19-5P
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (nucleotide sequence; vaccines comprising optimized multiple cytotoxic
        or helper T lymphocyte epitopes derived from hepatitis B virus pol, env
        and core proteins)
                                               139639-23-9, Tissue plasminogen
     9004-10-8, Insulin, biological studies
IT
     activator
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (signal sequence; vaccines comprising optimized multiple cytotoxic or
        helper T lymphocyte epitopes derived from hepatitis B virus pol, env
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        (unclaimed sequence; vaccines comprising optimized multiple cytotoxic
        or helper T lymphocyte epitopes derived from hepatitis B virus pol, env
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        (vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
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     RL: BSU (Biological study, unclassified); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (vaccines comprising optimized multiple cytotoxic or helper T
        lymphocyte epitopes derived from hepatitis B virus pol, env and core
        proteins)
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     RL: PRP (Properties)
        (unclaimed sequence; vaccines comprising optimized multiple cytotoxic
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        and core proteins)
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CN
     histidyl-L-seryl- (9CI) (CA INDEX NAME)
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Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

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ANSWER 5 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
     2004:101303 HCAPLUS
AN
      140:180120
DN
     Entered STN: 08 Feb 2004
ED
     Alternative reading frame (ncORF) antigenic determinants from viruses and
ΤI
      uses in vaccines
     Mattner, Frank; Schmidt, Walter; Habel, Andre
IN
      Intercell A.-G., Austria
PΑ
SO
      PCT Int. Appl., 220 pp.
      CODEN: PIXXD2
DT
      Patent
LΑ
      English
     ICM C12N015-33
IC
     ICS C07K014-005; C07K014-18; A61K039-12
      15-2 (Immunochemistry)
      Section cross-reference(s): 3, 10
FAN.CNT 7
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                                                                             DATE
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                          C12N015-33
 WO 2004011650
                  ICS
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                          A61K039/12; A61K039/145; A61K039/29B; C07K014/005;
 WO 2004011650
                  ECLA
                          C07K014/11; C07K014/18F4
     It is an object of the present invention to provide means for replacing or
AB
     improving existing or proposed vaccines against viral pathogens, especially
     human pathogens. A specific aim is to provide effective T cell epitopes
     against viral pathogens. The invention discloses polypeptides encoded by
     an alternative reading frame (non-coding open-reading frame (ncORF)) of a
     pathogenic virus, which polypeptides - start with a methionine amino acid
     residue, - comprise an antigenic determinant (epitope) and - comprise more
     than 7 amino acid residues and fragments of said polypeptides comprising
     more than 7 amino acids. T cell responses against alternatively encoded
     epitopes are detectable in patients suffering such infections. Such a
     polypeptide according to the present invention may be defined as an
     antigenic sequence outside the primarily (main) transcribed ORF of a given
     pathogenic virus. Alternatively encoded antigens from hepatitis C virus
     and human immunodeficiency virus are provided. Possible ncORF epitopes
     with superior immunization properties were identified for hepatitis C
     virus (HCV), human immunodeficiency virus (HIV) and human papilloma virus
     (HPV). The immunogenicity of HCV ncORF peptides was demonstrated on
     HLA-A-allele-transgenic mice and on HCV patient -derived cells.
     alternative reading frame antigen virus vaccine; vaccine HCV HIV HPV virus
ST
     epitope ncORF alternative
ΙT
     Epitopes
         (CTL-; alternative reading frame (ncORF) antigenic determinants from
        viruses and uses in vaccines)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (HLA, allele, CTL-epitope for; alternative reading frame (ncORF)
         antigenic determinants from viruses and uses in vaccines)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (HLA-A, alleles A0201, A1, A24, A3, A31, CTL-epitope for; alternative
        reading frame (ncORF) antigenic determinants from viruses and uses in
        vaccines)
ΙT
     Gene, animal
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (HLA-B, alleles B3501, B4403, B7, B8, CTL-epitope for; alternative
         reading frame (ncORF) antigenic determinants from viruses and uses in
         vaccines)
     Histocompatibility antigens
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (HLA-DP, alleles, CTL-epitopes for; alternative reading frame (ncORF)
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antiquenic determinants from viruses and uses in vaccines)
ΙT
    Gene, animal
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-DQ, alleles, CTL-epitope for; alternative reading frame (ncORF)
       antigenic determinants from viruses and uses in vaccines)
IT
    Gene, animal
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-DR, alleles, CTL-epitopes for; alternative reading frame (ncORF)
        antigenic determinants from viruses and uses in vaccines)
ΙT
    Antigens
    RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (T cell; alternative reading frame (ncORF) antigenic determinants from
        viruses and uses in vaccines)
IT
    Antiviral agents
    Cytomegalovirus
    Ebola virus
    Feline immunodeficiency virus
    Foot-and-mouth disease virus
    Hepatitis A virus
    Hepatitis B virus
    Hepatitis C virus
    Hepatitis E virus
    Hepatitis F virus
    Hepatitis GB virus C/G
    Hepatitis delta virus
    Human
    Human T-lymphotropic virus 1
    Human T-lymphotropic virus 2
    Human adenovirus
    Human herpesvirus
    Human herpesvirus 3
    Human herpesvirus 4
    Human immunodeficiency virus
    Human immunodeficiency virus 1
    Human papillomavirus
     Immunomodulators
     Influenza virus
    Measles virus
     Papillomavirus
    Parvovirus
     Rotavirus
     Simian immunodeficiency virus
        (alternative reading frame (ncORF) antigenic determinants from viruses
        and uses in vaccines)
IT
    Vaccines
        (antiviral; alternative reading frame (ncORF) antigenic determinants
        from viruses and uses in vaccines)
IT
    Drug delivery systems
        (carriers; alternative reading frame (ncORF) antigenic determinants
        from viruses and uses in vaccines)
IT
     Immunostimulants
        (co-application with; alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
IT
    Anti-inflammatory agents
    Antimicrobial agents
        (co-application; alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
IT
        (controlling; alternative reading frame (ncORF) antigenic determinants
        from viruses and uses in vaccines)
     T cell (lymphocyte)
IT
        (cytotoxic, CTL, epitope; alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
     Oligodeoxyribonucleotides
IT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
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(deoxyinosine- and/or deoxyuridine-containing, immunomodulating, conjugated
        to ncORF epitope; alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
     T cell (lymphocyte)
IT
        (helper cell, epitope; alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
     Cytokines
IT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (immunopotentiating, co-application; alternative reading frame (ncORF)
        antigenic determinants from viruses and uses in vaccines)
IT
        (of alternatively encoded antigens from hepatitis C virus and human
        immunodeficiency virus; alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
     Animal virus
TT
        (oncogenic; alternative reading frame (ncORF) antigenic determinants
        from viruses and uses in vaccines)
IT
     Gene
     RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological
     study); USES (Uses)
        (open reading frame, alternative, for complementary strand, non-coding
        open-reading frame (ncORF); alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
ΙT
     Virus
        (pathogenic; alternative reading frame (ncORF) antigenic determinants
        from viruses and uses in vaccines)
     Peptides, biological studies
TТ
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (polycationic, immunomodulating, conjugated to ncORF epitope;
        alternative reading frame (ncORF) antigenic determinants from viruses
        and uses in vaccines)
     Peptides, biological studies
тт
     Polymers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (polycationic, immunostimulatory; alternative reading frame (ncORF)
        antigenic determinants from viruses and uses in vaccines)
     9002-72-6, Growth hormone 170006-50-5, Cathelicidin
IT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (-derived peptide, immunostimulatory; alternative reading frame (ncORF)
        antigenic determinants from viruses and uses in vaccines)
     25212-18-4, L-Arginine, homopolymer
TΤ
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
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        viruses and uses in vaccines)
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     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
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        (amino acid sequence; alternative reading frame (ncORF) antigenic
        determinants from viruses and uses in vaccines)
     890-38-0, Deoxyinosine 951-78-0, Deoxyuridine
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        (immunomodulating oligodeoxyribonucleotide containing; alternative reading
        frame (ncORF) antigenic determinants from viruses and uses in vaccines)
IT
     25104-18-1, L-Lysine, homopolymer
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
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        determinants from viruses and uses in vaccines)
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IT 657367-96-9 657380-51-3

RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(amino acid sequence; alternative reading frame (ncORF) antigenic determinants from viruses and uses in vaccines)

RN 657367-96-9 HCAPLUS

CN L-Isoleucine, glycyl-L-prolyl-L-histidyl-L-threonyl-L-cysteinyl-L- α -aspartyl-L-alanylglycyl-L-threonyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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CN L-Isoleucine, L-alanyl-L-prolyl-L-histidyl-L-threonyl-L-methionyl-L-threonyl-L-phenylalanyl-L-phenylalanyl-L-seryl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B

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ANSWER 6 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
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     140:110124
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     Entered STN: 18 Jan 2004
ED
     Glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify anti-platelet
TI
     autoantibodies and inhibitors and for diagnosis and therapy of idiopathic
     thrombocytopenic purpura
IN
     Siegel, Donald L.
     The Trustees of the University of Pennsylvania, USA
PΑ
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     PCT Int. Appl., 232 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LΑ
     ICM G01N
IC
     15-3 (Immunochemistry)
CC
     Section cross-reference(s): 1, 3, 9, 63
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     WO 2004005890
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             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
         UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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CLASS
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PATENT NO.
WO 2004005890
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                        C07K016/00A; C07K016/28B14; G01N033/86
WO 2004005890 ECLA
                       C07K016/00A; C07K016/28B14; G01N033/86
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EP 1539236
                ECLA
    The present invention relates to novel methods of identifying and
    producing an anti-platelet autoantibody. More preferably, the invention
    relates to identification and production of a human monoclonal anti-platelet
     autoantibody. Addnl., the invention relates to methods for producing and
     identifying inhibitors of an anti-platelet autoantibody binding with a
     platelet, or a platelet component e.g. glycoproteins GPIa/IIa, GPIIb/IIIa
     and GPIb/IX. Moreover, the invention relates to methods for treating or
     alleviating a disease, disorder or condition mediated by an anti-platelet
     autoantibody specifically binding with a platelet, or a component thereof,
     such as, but not limited to, idiopathic thrombocytopenic purpura, among
     others.
     glycoprotein GPIaIIa GPIIbIIIa GPIbIX autoantibody platelet idiopathic
     thrombocytopenic purpura; phage display library antibody screening
     platelet adhesion aggregation glycoprotein
     RL: BSU (Biological study, unclassified); BUU (Biological use;
     unclassified); BIOL (Biological study); USES (Uses)
        (A, Staphylococcal; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
IT
     Glycoproteins
     RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
     DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
        (GP IX, complex with Ib and V; glycoproteins GPIa/IIa, GPIIb/IIIa and
        GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
     Glycoproteins
IT
     RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
     DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
        (GP V, complex with Ib and IX; glycoproteins GPIa/IIa, GPIIb/IIIa and
        GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
TT
     Glycoproteins
     RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
     DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
        (GPIb, complex with IX and V; glycoproteins GPIa/IIa, GPIIb/IIIa and
        GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
     Antibodies and Immunoglobulins
IT
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (IgG, immobilized; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
     Antibodies and Immunoglobulins
TT
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
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(IqG, monoclonal; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
    Antibodies and Immunoglobulins
TΤ
    RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (IgG; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
     Platelet (blood)
IT
        (adhesion, inhibitors; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX
        to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
     Antibodies and Immunoglobulins
IT
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (autoantibodies, monoclonal; glycoproteins GPIa/IIa, GPIIb/IIIa and
        GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
     Antibodies and Immunoglobulins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (autoantibodies; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
IT
     Fibronectins
     RL: ARU (Analytical role, unclassified); BSU (Biological study,
     unclassified); ANST (Analytical study); BIOL (Biological study)
        (binding inhibition; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
     Antibodies and Immunoglobulins
IT
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (fragments; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
IT
     Anticoagulants
     Autoimmune disease
     B cell (lymphocyte)
     Blood coagulation
     DNA sequences
     Drug screening
     Human
     Immunotherapy
     Mammalia
     Medical goods
     Molecular cloning
     Panning
     Phage display library
     Platelet (blood)
     Platelet aggregation inhibitors
     Protein sequences
     Test kits
     cDNA sequences
        (qlycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
     Gene, animal
TТ
     RL: ARU (Analytical role, unclassified); BPN (Biosynthetic preparation);
     BSU (Biological study, unclassified); ANST (Analytical study); BIOL
      (Biological study); PREP (Preparation)
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(qlycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
IT
    Nucleic acids
     RL: ARU (Analytical role, unclassified); BSU (Biological study,
     unclassified); ANST (Analytical study); BIOL (Biological study)
        (qlycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
IT
     Antibodies and Immunoglobulins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (heavy chain; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
IT
     Purpura (disease)
        (idiopathic thrombocytopenic; glycoproteins GPIa/IIa, GPIIb/IIIa and
        GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
TT
     Diagnosis
        (immunodiagnosis; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
     Drug delivery systems
TT
        (immunotoxins; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
     Peptides, biological studies
TΤ
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (inhibitor; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
TΤ
     Thrombus
        (inhibitors; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
     Antibodies and Immunoglobulins
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (library; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
     Antibodies and Immunoglobulins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (light chain; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
IT
     Epitopes
        (mapping; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
IT
     Antibodies and Immunoglobulins
     RL: ANT (Analyte); \overline{	ext{BSU}} (Biological study, unclassified); \overline{	ext{DGN}} (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (monoclonal; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
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(platelet, inhibitors; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX

Adhesion, biological

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to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
IT
    Staphylococcus
        (protein A; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
TT
    Receptors
    RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
    DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
        (von Willebrand factor; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX
        to identify anti-platelet autoantibodies and inhibitors and for
        diagnosis and therapy of idiopathic thrombocytopenic purpura)
    Integrins
    RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
    DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
        (\alpha IIb\beta 3; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
IT
    Integrins
    RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
    DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
        (\alpha 2\beta 1; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
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     BSU (Biological study, unclassified); DGN (Diagnostic use); PRP
     (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (amino acid sequence; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to
        identify anti-platelet autoantibodies and inhibitors and for diagnosis
        and therapy of idiopathic thrombocytopenic purpura)
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     unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
        (glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify
        anti-platelet autoantibodies and inhibitors and for diagnosis and
        therapy of idiopathic thrombocytopenic purpura)
     109319-16-6
IT
     RL: ARU (Analytical role, unclassified); BSU (Biological study,
     unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
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(multimer; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for diagnosis and therapy of idiopathic thrombocytopenic purpura)

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(nucleotide sequence; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for diagnosis and therapy of idiopathic thrombocytopenic purpura)

IT 173319-74-9 311337-02-7

RL: PRP (Properties)

(unclaimed sequence; glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for diagnosis and therapy of idiopathic thrombocytopenic purpura)

IT 647825-82-9

IT

RL: ARU (Analytical role, unclassified); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)

(glycoproteins GPIa/IIa, GPIIb/IIIa and GPIb/IX to identify anti-platelet autoantibodies and inhibitors and for diagnosis and therapy of idiopathic thrombocytopenic purpura)

RN 647825-82-9 HCAPLUS

CN L-Cysteine, L-cysteinyl-L-valyl-L-threonyl-L-seryl-L-lysyl-L-prolyl-L-histidyl-L-threonyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 2-A

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ANSWER 7 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
     2004:3693 HCAPLUS
AN
     140:72162
     Entered STN: 04 Jan 2004
ED
     Cloning and cDNA and deduced amino acid sequences of 50 human secreted
TI
     Moore, Paul A.; Ruben, Steven M.; Lafleur, David W.; Shi, Yanggu; Rosen,
TN
     Craiq A.; Olsen, Henrik S.; Ebner, Reinhard; Brewer, Laurie A.
     Human Genome Sciences, Inc., USA
PΑ
     U.S. Pat. Appl. Publ., 383 pp., Cont.-in-part of U.S. Ser. No. 722,329.
SO
     CODEN: USXXCO
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LA
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     ICS C07K014-00; C07K017-00
INCL 530395000
     3-3 (Biochemical Genetics)
     Section cross-reference(s): 6, 13, 63
FAN.CNT 43
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                                                                           DATE
     PATENT NO.
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 US 2004002591
                           C07K014-00; C07K017-00
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                   NCL
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                           C07K014/47
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 WO 9911293
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AB The present invention relates to 50 novel human secreted proteins and
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isolated nucleic acids containing the coding regions of the genes encoding
such proteins. Tissue distribution, sequence homologies, and preferred
epitope sites are provided for the secreted proteins, as well as
chromosomal mapping of some of the genes. Also provided are vectors, host
cells, antibodies, and recombinant methods for producing human secreted
proteins in bacterial, insect, and mammalian cells. The invention further
relates to diagnostic and therapeutic methods useful for diagnosing and
treating disorders related to these novel human secreted proteins.
High-throughput screening assays are also provided for various putative
activities of the secreted proteins.
secretory protein cDNA sequence human
High throughput screening
   (assays; cloning and cDNA and deduced amino acid sequences of 50 human
   secreted proteins)
Epitopes
Human
Molecular cloning
Protein sequences
cDNA sequences
   (cloning and cDNA and deduced amino acid sequences of 50 human secreted
   proteins)
Antibodies and Immunoglobulins
RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
   (cloning and cDNA and deduced amino acid sequences of 50 human secreted
   proteins)
Gene, animal
RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
(Biological study)
   (cloning and cDNA and deduced amino acid sequences of 50 human secreted
   proteins)
Disease, animal
   (diagnosis and treatment of; cloning and cDNA and deduced amino acid
   sequences of 50 human secreted proteins)
Animal tissue
   (expression profiles; cloning and cDNA and deduced amino acid sequences
   of 50 human secreted proteins)
Genetic mapping
   (gene location on human chromosome; cloning and cDNA and deduced amino
   acid sequences of 50 human secreted proteins)
Chromosome
   (human, gene location on; cloning and cDNA and deduced amino acid
   sequences of 50 human secreted proteins)
   (mol., by mutation detection or changes in expression; cloning and cDNA
   and deduced amino acid sequences of 50 human secreted proteins)
Proteins
RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL
(Biological study); PREP (Preparation); USES (Uses)
   (secretory; cloning and cDNA and deduced amino acid sequences of 50
   human secreted proteins)
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RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL
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(Biological study); PREP (Preparation); USES (Uses)

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(amino acid sequence; cloning and cDNA and deduced amino acid sequences
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Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

PAGE 1-C

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PAGE 1-D

PAGE 1-E

PATENT NO.

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AN
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ED
     Entered STN: 23 Oct 2003
     Human genome-derived single exon nucleic acid probes useful for analysis
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     of gene expression in human tissues
     Penn, Sharron Gaynor; Rank, David Russell; Hanzel, David Kagen
IN
PΑ
     USA
     U.S. Pat. Appl. Publ., 80 pp.
SO
     CODEN: USXXCO
DT
     Patent
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     English
     C12Q001-68; C07H021-04
IC
INCL 435006000; 536024300
     3-3 (Biochemical Genetics)
     Section cross-reference(s): 6, 13
FAN.CNT 9
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CLASS
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CLASS PATENT FAMILY CLASSIFICATION CODES

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                 NCL
                        435/006.000; 536/024.300
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    Methods and apparatus for predicting, confirming and displaying functional
    regions from genomic sequence data are used to identify unique human
    genome-derived single exon probes useful for gene expression anal.,
    particularly gene expression anal. by microarray. Human BAC sequences
    totaling .apprx.350 Mb of sequence (.apprx.10% of the human genome) were
    analyzed for exons using four sep. gene finding programs (GRAIL uses a
    neural network, GENEFINDER uses a hidden Markoff model, DICTION operates
    according to a different heuristic, and GENSCAN) and Mouse comparative
    genomics as a fifth gene prediction method. The exons were PCR amplified from genomic DNA, verified on agarose gels, and sequenced using universal
    primers to validate the identity of the amplicon to be spotted in
    microarrays. Thus, expression, homol., and functional information are
    provided for 13,700 unique genome-derived single exon probes are expressed
    at significant levels in one or more of 8 tested tissues: human brain,
    heart, fetal and adult liver, placenta, lung, bone marrow, and HeLa cells.
    The probes lack prokaryotic and bacteriophage vector sequences, as well as
    lacking homopolymeric stretches of A or T. Also presented are
    genome-derived single exon microarrays that include such probes, peptides
    encoded by the exons, and antibodies thereto. In addition, methods are
    provided for identifying exons in a eukaryotic genome, and for assigning
     exons to a single gene. [This abstract record is one of nine records for
     this document necessitated by the large number of index entries required to
     fully index the document and publication system constraints.].
     genome exon probe microarray gene expression human lung; sequence genome
     exon probe human lung
IT
    Liver
        (adult and fetal; human genome-derived single exon nucleic acid probes
        useful for anal. of gene expression in human tissues)
IT
     Information systems
        (computerized, computer-readable storage medium; human genome-derived
        single exon nucleic acid probes useful for anal. of gene expression in
        human tissues)
IT
     Genetic element
     RL: ANT (Analyte); BUU (Biological use, unclassified); PRP (Properties);
     ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (exon; human genome-derived single exon nucleic acid probes useful for
        anal. of gene expression in human tissues)
     Bone marrow
IT
     Brain
     DNA microarray technology
     DNA sequences
     Gene expression profiles, animal
     Genetic vectors
     HeLa cell
     Heart
     Human
     Lung
     Molecular cloning
     Placenta
     Protein sequences
        (human genome-derived single exon nucleic acid probes useful for anal.
        of gene expression in human tissues)
     Gene, animal
IT
     RL: ANT (Analyte); BSU (Biological study, unclassified); PRP (Properties);
     ANST (Analytical study); BIOL (Biological study)
        (human genome-derived single exon nucleic acid probes useful for anal.
        of gene expression in human tissues)
     Peptides, biological studies
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     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); BIOL (Biological study); USES (Uses)
        (human genome-derived single exon nucleic acid probes useful for anal.
        of gene expression in human tissues)
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IT
      Antibodies and Immunoglobulins
      RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
      (Uses)
         (human genome-derived single exon nucleic acid probes useful for anal.
         of gene expression in human tissues)
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      Probes (nucleic acid)
      RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological
      study); USES (Uses)
         (human genome-derived single exon nucleic acid probes useful for anal.
         of gene expression in human tissues)
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         acid probes useful for anal. of gene expression in human tissues)
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      Lung
         (toxicity; human genome-derived single exon nucleic acid probes useful
         for anal. of gene expression in human tissues)
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BUU (Biological use, unclassified); PRP (Properties); ANST (Analytical
study); BIOL (Biological study); USES (Uses)
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BUU (Biological use, unclassified); PRP (Properties); ANST (Analytical
study); BIOL (Biological study); USES (Uses)
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        (amino acid sequence; human genome-derived single exon nucleic acid
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     study); BIOL (Biological study); USES (Uses)
        (amino acid sequence; human genome-derived single exon nucleic acid
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BUU (Biological use, unclassified); PRP (Properties); ANST (Analytical
study); BIOL (Biological study); USES (Uses)
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BUU (Biological use, unclassified); PRP (Properties); ANST (Analytical
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610776-60-8
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610776-60-8 HCAPLUS
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IT

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CN

alanyl-L- α -aspartyl-L-leucyl-L-leucyl- (9CI) (CA INDEX NAME) Absolute stereochemistry.

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ANSWER 9 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

2003:719271 HCAPLUS AN

DN 139:265740

ED Entered STN: 14 Sep 2003

KDR and VEGF/KDR binding peptides and their use in diagnosis and therapy ΤI

Sato, Aaron K.; Sexton, Daniel J.; Ladner, Robert C.; Dransfield, Daniel T.; Swenson, Rolf E.; Marinelli, Edmund R.; Ramalingam, Kondareddiar; IN Nunn, Adrian D.; Von Wronski, Mathew A.; Shrivastava, Ajay; Pochon, Sibylle; Bussat, Philippe; Arbogast, Christophe; Pillai, Radhakrishna; Fan, Hong; Linder, Karen E.; Song, Bo; Nanjappan, Palaniappa

Dyax Corp., USA; Bracco International B.V.; et al. PA

so PCT Int. Appl., 350 pp.

CODEN: PIXXD2

DTPatent

LА English

ICM A61K IC

63-5 (Pharmaceuticals) CC

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Section cross-reference(s): 1, 8, 9
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     The present invention relates to polypeptides useful for detecting and
AB
     targeting primary receptors on endothelial cells for VEGF, i.e., VEGF
      receptor 2, also known as kinase domain region (KDR) and fetal liver
      kinase-1 (Flk-1), and for imaging and targeting complexes formed by VEGF
     and KDR. The involvement of VEGF and KDR in angiogenesis makes the
     VEGF/KDR and KDR binding polypeptides of the present invention
     particularly useful for imaging important sites of angiogenesis, e. g.,
     neoplastic tumors, for targeting substances, e.g., therapeutics, including radiotherapeutics, to such sites, and for treating certain disease states,
      including those associated with inappropriate angiogenesis. Disclosed are
      synthetic, isolated polypeptides capable of binding KDR or VEGF/KDR
      complex with high affinity (e.g., having a KD<1 \muM).
     kinase domain region VEGF targeting peptide diagnosis therapy; imaging
ST
     radiotherapy tumor VEGF receptor target
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     AIDS (disease)
     Anti-AIDS agents
     Antimalarials
     Antitumor agents
     Antiviral agents
      Chelating agents
     Diagnosis
     Drug targets
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Fluorescent indicators
     Imaging agents
     Malaria
     Phage display library
     Radiopharmaceuticals
     Radiotherapy
     Scintigraphic agents
     Simian hemorrhagic fever virus
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     Imaging agents
        (NMR contrast; KDR and VEGF/KDR binding peptides and their use in
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        diagnosis and therapy)
     Drug delivery systems
IT
        (liposomes; KDR and VEGF/KDR binding peptides and their use in
        diagnosis and therapy)
IT
     Bubbles
        (microbubbles, ultrasound contrast agents; KDR and VEGF/KDR binding
        peptides and their use in diagnosis and therapy)
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150977-45-0, Kdr

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biological studies
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studies 14158-31-7D, Iodine 125, compds., biological studies
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69146-59-4D, MECAM, conjugates 71353-07-6D, conjugates
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DOTMA, conjugates 149440-35-7D, DO3A, conjugates
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Neutravidin, conjugates 325126-08-7D, conjugates
RL: DGN (Diagnostic use); THU (Therapeutic use); BIOL (Biological study);
USES (Uses)
   (KDR and VEGF/KDR binding peptides and their use in diagnosis and
   therapy)
1190-93-8, S-Acetylthioglycolic acid
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599211-93-5D, resin-bound 599211-94-6D, resin-bound
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   (KDR and VEGF/KDR binding peptides and their use in diagnosis and
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   (KDR and VEGF/KDR binding peptides and their use in diagnosis and
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13981-59-4D, Tin 117, compds., biological studies
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USES (Uses)
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RL: PRP (Properties)

(unclaimed sequence; kDR and VEGF/KDR binding peptides and their use in diagnosis and therapy)

IT 599210-01-2P

CN

RL: DGN (Diagnostic use); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (KDR and VEGF/KDR binding peptides and their use in diagnosis and therapy)

RN 599210-01-2 HCAPLUS

L-Glutamic acid, glycyl-L-histidyl-L-prolyl-L-cysteinyl-L-lysylglycyl-L-methionyl-L-leucyl-L-prolyl-L-histidyl-L-threonyl-L-cysteinyl-L-tryptophyl-L-tyrosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

HO2C
$$\frac{CO_2H}{H}$$
 $\frac{S}{H}$ $\frac{S}{H}$ $\frac{H}{S}$ $\frac{H}{H}$ $\frac{S}{I-Bu}$

PAGE 1-B

L24 ANSWER 10 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:221865 HCAPLUS

DN 138:250714

ED Entered STN: 21 Mar 2003

TI Selection and production of laccase- and other phenol oxidizing enzyme-binding peptide complexes for detergents and carotenoid stain removal

IN Aehle, Wolfgang; Baldwin, Toby L.; Janssen, Giselle G.; Murray, Christopher J.; Van Gastel, Franciscus J. C.; Wang, Huaming; Winetzky, Deborah S.

PA Genencor International, Inc., USA

SO PCT Int. Appl., 50 pp. CODEN: PIXXD2

DT Patent

LA English

IC ICM C12S009-00

CC 7-3 (Enzymes)

Section cross-reference(s): 3, 16, 46

FAN.CNT 1

FAN.CNT 1 PATENT NO.		KIND	DATE	APPLICATION NO.	DATE
ΡI	WO 2003023067	A1	20030320	WO 2002-US27993	20020903 <
•	WO 2003023067	C2	20040521		

Search done by Noble Jarrell

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CLASS
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                        C11D003/386H; C12N009/02H3B2
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                        510/392.000; 510/530.000; 435/183.000
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 US 2005042684
                        435/007.100; 435/069.100; 435/189.000; 435/320.100;
                        435/325.000; 530/388.260; 536/023.200
                        C11D003/386H; C12N009/02H3B2
                 ECLA
AB
     The present application relates to peptides which bind to a target stain,
     phenol oxidizing enzyme-binding peptide complexes wherein the binding
     peptide is attached to the C-terminus of the phenol oxidizing enzyme or is
     inserted or substituted into the phenol oxidizing enzyme. In a preferred
     embodiment the phenol oxidizing enzyme is a laccase, specifically Stachybotrys phenol oxidase B and variants thereof. The invention
     provides expression vectors comprising the phenol oxidizing enzyme-binding
     peptide complex as well as host cells comprising the vectors. The phenol
     oxidizing enzyme-binding peptide complex may be used in enzymic compns.,
     particularly detergent compns. to specifically target stains and
     particularly to carotenoid compound stains. The invention pertain to a
     method of using a binding peptide to target a stain on a textile.
st
     phenol oxidizing enzyme peptide complex detergent carotenoid stain
     removal; laccase peptide complex detergent carotenoid stain removal;
     Stachybotrys phenol oxidase B peptide complex detergent carotenoid stain
IT
     Protein sequences
        (Stachybotrys phenol oxidase B; selection and production of laccase- and
        other phenol oxidizing enzyme-binding peptide complexes for detergents
        and carotenoid stain removal)
IT
     Aspergillus niger
        (cloning host; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
        carotenoid stain removal)
IT
     Detergents
        (enzyme-containing; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
        carotenoid stain removal)
IT
     Genetic vectors
        (for enzyme-peptide complexes; selection and production of laccase- and
        other phenol oxidizing enzyme-binding peptide complexes for detergents
        and carotenoid stain removal)
IT
     Molecular cloning
        (of enzyme-peptide complexes; selection and production of laccase- and
        other phenol oxidizing enzyme-binding peptide complexes for detergents
        and carotenoid stain removal)
IT
     Repeat motifs (protein)
        (of peptide; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
        carotenoid stain removal)
IT
     Molecular association
        (peptide-enzyme and peptide-carotenoid; selection and production of
        laccase- and other phenol oxidizing enzyme-binding peptide complexes
        for detergents and carotenoid stain removal)
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IT

Stachybotrys

```
(phenol oxidase B; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
        carotenoid stain removal)
     Enzymes, biological studies
     RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); CAT (Catalyst use); NUU (Other use, unclassified); PRP (Properties); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (phenol oxidizing; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
        carotenoid stain removal)
     Fermentation
IT
        (protein, of enzyme-peptide complexes; selection and production of laccase-
        and other phenol oxidizing enzyme-binding peptide complexes for
        detergents and carotenoid stain removal)
IT
     Bleaching
     Bleaching agents
     Coloring materials
     Genetic engineering
     Phage display library
     Stains, biological
     Surfactants
        (selection and production of laccase- and other phenol oxidizing
        enzyme-binding peptide complexes for detergents and carotenoid stain
     Peptides, biological studies
IT
     RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); NUU (Other use, unclassified); PRP
     (Properties); BIOL (Biological study); PREP (Preparation); USES (Uses)
         (selection and production of laccase- and other phenol oxidizing
        enzyme-binding peptide complexes for detergents and carotenoid stain
     Carotenes, biological studies
ΙT
     RL: BSU (Biological study, unclassified); REM (Removal or disposal); BIOL
     (Biological study); PROC (Process)
         (selection and production of laccase- and other phenol oxidizing
        enzyme-binding peptide complexes for detergents and carotenoid stain
        removal)
     Detergents
IT
         (stain removers; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
        carotenoid stain removal)
     Lycopersicon esculentum
IT
     Textiles
         (stain; selection and production of laccase- and other phenol oxidizing
         enzyme-binding peptide complexes for detergents and carotenoid stain
        removal)
     9002-10-2P, Phenol oxidase
IT
     RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); CAT (Catalyst use); NUU (Other use, unclassified); PRP (Properties); BIOL (Biological study); PREP
      (Preparation); USES (Uses)
         (B; selection and production of laccase- and other phenol oxidizing
        enzyme-binding peptide complexes for detergents and carotenoid stain
         removal)
     502774-31-4DP, variants are claimed
IT
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      (Biological study, unclassified); CAT (Catalyst use); NUU (Other use,
     unclassified); PRP (Properties); BIOL (Biological study); PREP
      (Preparation); USES (Uses)
         (amino acid sequence; selection and production of laccase- and other phenol
         oxidizing enzyme-binding peptide complexes for detergents and
         carotenoid stain removal)
     52-90-4, L-Cysteine, biological studies
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     RL: BSU (Biological study, unclassified); BIOL (Biological study)
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(peptide containing; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
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     (Biological study)
        (protein motif; selection and production of laccase- and other phenol
        oxidizing enzyme-binding peptide complexes for detergents and
        carotenoid stain removal)
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     80498-15-3P, Laccase
     RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); CAT (Catalyst use); NUU (Other use, unclassified); PRP (Properties); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (selection and production of laccase- and other phenol oxidizing
        enzyme-binding peptide complexes for detergents and carotenoid stain
        removal)
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        (selection and production of laccase- and other phenol oxidizing
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        (unclaimed nucleotide sequence; selection and production of laccase- and
        other phenol oxidizing enzyme-binding peptide complexes for detergents
        and carotenoid stain removal)
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RL: PRP (Properties)

(unclaimed sequence; selection and production of laccase- and other phenol oxidizing enzyme-binding peptide complexes for detergents and carotenoid stain removal)

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

(1) Wang; US 6168936 B1 2001 HCAPLUS

IT 502169-28-0

RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(protein motif; selection and production of laccase- and other phenol oxidizing enzyme-binding peptide complexes for detergents and carotenoid stain removal)

RN 502169-28-0 HCAPLUS

CN L-Methionine, L-prolyl-L-histidyl-L-seryl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 502166-83-8P 502167-78-4P 502167-93-3P

RL: BMF (Bioindustrial manufacture); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); NUU (Other use, unclassified); PRP (Properties); BIOL (Biological study); PREP (Preparation); USES (Uses) (selection and production of laccase- and other phenol oxidizing enzyme-binding peptide complexes for detergents and carotenoid stain removal)

RN 502166-83-8 HCAPLUS

CN L-Methionine, L-isoleucyl-L-asparaginyl-L-threonyl-L-prolyl-L-histidyl-L-seryl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN

CN L-Proline, L-seryl-L-prolyl-L-histidyl-L-seryl-L-methionyl-L-leucyl-L-glutaminyl-L-asparaginyl-L-prolyl-L-serylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A.

PAGE 1-B

RN 502167-93-3 HCAPLUS

CN L-Tryptophan, L-valyl-L-alanyl-L-seryl-L-alanyl-L-asparaginyl-L-prolyl-L-histidyl-L-seryl-L-methionyl-L-threonyl-L-seryl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

PAGE 2-B

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L24 ANSWER 11 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:889383 HCAPLUS

137:381699

DN

ED Entered STN: 22 Nov 2002

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Crystal structure of Drosophila \alpha-mannosidase II and swainsonine
     complexes, its use for identifying mannosidase II activity-modulating
     ligands and therapeutic applications
IN
     Rose, David Richard; Kuntz, Douglas Arthur; Van Den Elsen, Jean Maria
    Hubertus
PΑ
    Can.
    U.S. Pat. Appl. Publ., 429 pp.
    CODEN: USXXCO
DT
    Patent
LΑ
    English
    ICM C12N009-24
TC
     ICS A61K038-47; G06F019-00; G01N033-48; G01N033-50
INCL 424094610; 435200000; 702019000
     7-5 (Enzymes)
    Section cross-reference(s): 1, 9, 63, 75
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                                           APPLICATION NO.
                                                                   DATE
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CLASS
                CLASS PATENT FAMILY CLASSIFICATION CODES
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                        C12N009-24
 US 2002172670
                ICM
                       A61K038-47; G06F019-00; G01N033-48; G01N033-50
                 ICS
                       424094610; 435200000; 702019000
                 INCL
                        424/094.610; 435/200.000; 702/019.000
 US 2002172670
                NCL
                        C12Q001/34; G01N033/68A
                 ECLA
    The present invention relates to a crystal comprising a mannosidase II
AR
     ligand-binding domain. In particular the present invention relates to a
     crystal comprising Drosophila Golgi \alpha\text{-mannosidase} II (EC 3.2.1.114)
     (with and without swainsonine), and its use to generate models for
     elucidating the structure of other polypeptides and for better identifying
     ligands capable of modulating mannosidase II activity. These ligands may
     have therapeutic and prophylactic potential.
    Drosophila mannosidase II swainsonine crystal structure ligand therapeutic
st
    Protein motifs
IT
        (Iq-like domain; crystal structure of Drosophila \alpha-mannosidase II
        and swainsonine complexes, its use for identifying mannosidase II
        activity-modulating ligands and therapeutic applications)
     Oligosaccharides, biological studies
IT
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (active site bound; crystal structure of Drosophila \alpha\text{-mannosidase}
        II and swainsonine complexes, its use for identifying mannosidase II
        activity-modulating ligands and therapeutic applications)
IT
     Enzyme functional sites
        (active; crystal structure of Drosophila \alpha\text{-mannosidase} II and
        swainsonine complexes, its use for identifying mannosidase II
        activity-modulating ligands and therapeutic applications)
TT
     Crystal growth
     DNA sequences
     Disulfide group
     Drosophila
     Drug screening
     Molecular modeling
     Protein motifs
     Protein sequences
     \alpha-Helix
        (crystal structure of Drosophila \alpha-mannosidase II and swainsonine
        complexes, its use for identifying mannosidase II activity-modulating
        ligands and therapeutic applications)
IT
     Ligands
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RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
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        (crystal structure of Drosophila \alpha-mannosidase II and swainsonine
        complexes, its use for identifying mannosidase II activity-modulating
        ligands and therapeutic applications)
     Information systems
IT
        (data, crystal structure data storage and mol. modeling; crystal
        structure of Drosophila \alpha-mannosidase II and swainsonine
        complexes, its use for identifying mannosidase II activity-modulating
        ligands and therapeutic applications)
IT 
     Protein motifs
        (glycosylation site; crystal structure of Drosophila
        \alpha-mannosidase II and swainsonine complexes, its use for
        identifying mannosidase II activity-modulating ligands and therapeutic
        applications)
IT
     Enzyme functional sites
       (ligand-binding; crystal structure of Drosophila \alpha-mannosidase II
        and swainsonine complexes, its use for identifying mannosidase II
        activity-modulating ligands and therapeutic applications)
IT
        (ligand-containing; crystal structure of Drosophila \alpha-mannosidase II
        and swainsonine complexes, its use for identifying mannosidase II
        activity-modulating ligands and therapeutic applications)
IT
     Golgi apparatus
        (mannosidase II from; crystal structure of Drosophila
        \alpha-mannosidase II and swainsonine complexes, its use for
        identifying mannosidase II activity-modulating ligands and therapeutic
        applications)
     Disease, animal
IT
        (mannosidase II-associated; crystal structure of Drosophila
        \alpha-mannosidase II and swainsonine complexes, its use for
        identifying mannosidase II activity-modulating ligands and therapeutic
        applications)
TT
     Enzyme functional sites
        (metal-binding, zinc-binding; crystal structure of Drosophila
        α-mannosidase II and swainsonine complexes, its use for
        identifying mannosidase II activity-modulating ligands and therapeutic
        applications)
TT
     Crystal structure
     Molecular structure
        (of Drosophila \alpha-mannosidase II and swainsonine complexes)
     Molecular association
IT
        (of ligand with mannosidase II; crystal structure of Drosophila
        \alpha-mannosidase II and swainsonine complexes, its use for
        identifying mannosidase II activity-modulating ligands and therapeutic
        applications)
     Molecular cloning
TT
        (of mannosidase II; crystal structure of Drosophila \alpha-mannosidase
        II and swainsonine complexes, its use for identifying mannosidase II
        activity-modulating ligands and therapeutic applications)
IT
     Coordination sphere
        (of zinc ion; crystal structure of Drosophila \alpha-mannosidase II
        and swainsonine complexes, its use for identifying mannosidase II
        activity-modulating ligands and therapeutic applications)
TT
     Plasmid vectors
        (pCopBlast, mannosidase II cloning; crystal structure of Drosophila
        \alpha\text{-mannosidase} II and swainsonine complexes, its use for
        identifying mannosidase II activity-modulating ligands and therapeutic
        applications)
IT
     Secondary structure
     Tertiary structure
        (protein, determination of secondary and tertiary structures using mannosidase
        II crystal structure; crystal structure of Drosophila
        α-mannosidase II and swainsonine complexes)
ΙT
     Conformation
        (protein; crystal structure of Drosophila \alpha\text{-mannosidase} II and
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swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

TT Information systems

(storage, crystal structure data storage and mol. modeling; crystal structure of Drosophila α -mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

TT Computer application

(to crystal structure data storage and mol. modeling; crystal structure of Drosophila α -mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

IT Mammalia

(treatment of disease of; crystal structure of Drosophila $\dot{\alpha}$ -mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

ΙT Conformation

(β-strand; crystal structure of Drosophila α-mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

·IT 70-47-3, L-Asparagine, biological studies

RL: BSU (Biological study, unclassified); BIOL (Biological study) (-194, N-glycosylation site; crystal structure of Drosophila α -mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

63-91-2, L-Phenylalanine, biological studies TT

RL: BSU (Biological study, unclassified); BIOL (Biological study) (-206, ligand-binding domain; crystal structure of Drosophila $\alpha\text{-mannosidase II}$ and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

60-18-4, L-Tyrosine, biological studies TT

RL: BSU (Biological study, unclassified); BIOL (Biological study) (-269, -727, ligand-binding domain; crystal structure of Drosophila α-mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

IT

52-90-4, L-Cysteine, biological studies RL: BSU (Biological study, unclassified); BIOL (Biological study) (-31 and -1032, -275-and 282, -283 and -297, disulfide bond; crystal structure of Drosophila α -mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

56-84-8, L-Aspartic acid, biological studies ΙT

RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(-90, -92, -204, -341, -472, ligand-binding domain; crystal structure of Drosophila α -mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

71-00-1, L-Histidine, biological studies IT

RL: BSU (Biological study, unclassified); BIOL (Biological study) (-90, -92, -471, ligand-binding domain; crystal structure of Drosophila lpha-mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

73-22-3, L-Tryptophan, biological studies TT

RL: BSU (Biological study, unclassified); BIOL (Biological study) (-95, ligand-binding domain; crystal structure of Drosophila $\alpha\text{-mannosidase II}$ and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

IT 77-86-1, Tris

RL: BSU (Biological study, unclassified); PRP (Properties); BIOL

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(Biological study)
             (active site bound; crystal structure of Drosophila \alpha-mannosidase
             II and swainsonine complexes, its use for identifying mannosidase II
             activity-modulating ligands and therapeutic applications)
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             (amino acid sequence; crystal structure of Drosophila
             \alpha-mannosidase II and swainsonine complexes, its use for
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             applications)
        7512-17-6, N-Acetyl-D-glucosamine
IT
        RL: BSU (Biological study, unclassified); BIOL (Biological study)
             (at N-glycosylation site; crystal structure of Drosophila
             lpha-mannosidase II and swainsonine complexes, its use for
             identifying mannosidase II activity-modulating ligands and therapeutic
             applications)
        7440-66-6, Zinc, biological studies
IT
        RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
        (Biological study)
              (at mannosidase II active site; crystal structure of Drosophila
             lpha-mannosidase II and swainsonine complexes, its use for
             identifying mannosidase II activity-modulating ligands and therapeutic
             applications)
        72741-87-8DP, Swainsonine, \alpha-mannosidase II complexes
IT
        82047-77-6DP, \alpha-Mannosidase II, swainsonine complexes
        RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
        BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological
        study); PREP (Preparation); USES (Uses)
              (crystal structure of Drosophila lpha-mannosidase II and swainsonine
             complexes, its use for identifying mannosidase II activity-modulating
             ligands and therapeutic applications)
        82047-77-6P, \alpha-Mannosidase II
TΤ
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             ligands and therapeutic applications)
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        study); PREP (Preparation); USES (Uses)
              (nucleotide sequence; crystal structure of Drosophila
             \alpha-mannosidase II and swainsonine complexes, its use for
             identifying mannosidase II activity-modulating ligands and therapeutic
             applications)
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             activity-modulating ligands and therapeutic applications)
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        L-Arginine, L-histidyl-L-methionyl-L-methionyl-L-prolyl-L-phenylalanyl-L-
CN
        tyrosyl-L-seryl-L-tyrosyl-L-\alpha-aspartyl-L-prolyl-L-histidyl-L-
        threonyl-L-cysteinylglycyl-L-prolyl-L-\alpha-aspartyl-L-prolyl-L-lysyl-L-aspartyl-L-prolyl-L-lysyl-L-aspartyl-L-prolyl-L-lysyl-L-aspartyl-L-prolyl-L-lysyl-L-aspartyl-L-prolyl-L-lysyl-L-aspartyl-L-prolyl-L-lysyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-prolyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl-L-aspartyl
        valyl-L-isoleucyl-L-cysteinyl-L-glutaminyl-L-phenylalanyl-L-\alpha- •
```

aspartyl-L-phenylalanyl-L-lysyl- (9CI) (CA INDEX NAME)
Absolute stereochemistry.

PAGE 1-B

PAGE 1-C

$$H_2N$$
 S
 H
 S
 H

PAGE 2-A

$$\begin{array}{c} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\$$

ANSWER 12 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN L24

2002:857442 HCAPLUS AN

137:321377 DN

Entered STN: 12 Nov 2002 ED

Human nucleic acids and corresponding proteins useful in the detection and ΤI treatment of various cancers

Jakobovits, Aya; Challita-Eid, Pia M.; Faris, Mary; Ge, Wangmao; Hubert, IN Rene S.; Morrison, Karen; Morrison, Robert Kendall; Raitano, Arthur B.

Agensys, Inc., USA PA

so PCT Int. Appl., 1021 pp.

CODEN: PIXXD2

DTPatent

LА English

IC C12Q

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 1, 6, 9, 14

FAN	.CNT 25				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	WO 2002083921	A2	20021024	WO 2002-XB11654	20020410 <

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     US 2003199470
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                                                 US 2002-280340
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     US 2003207835
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CLASS
                   CLASS PATENT FAMILY CLASSIFICATION CODES
PATENT NO.
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WO 2002083921
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US 2003199470
                   NCL
                           435/007.230
                           514/044.000; 424/146.100; 435/006.000; 435/007.230
                 NCL
 US 2003207835
     Eighteen genes and their resp. encoded proteins, and variants thereof, are
     described wherein the gene exhibits restricted expression in normal adult
     tissue and is overexpressed in various cancers. Suppression subtractive
     hybridization (SSH) is used to identify cDNAs corresponding to genes that
     are differentially expressed in cancer; PCR amplification, cloning, and
     sequencing of gene fragments from SSH yield the full-length cDNAs.
     Consequently, the gene products provide diagnostic, prognostic,
     prophylactic, and/or therapeutic targets for cancer. The genes or
     fragment thereof, their encoded proteins, or variants or fragments
     thereof, can be used to elicit a humoral or cellular immune response;
     antibodies or T cells reactive with the gene products can be used in
     active or passive immunization. [This abstract record is one of 16 records
     for this document necessitated by the large number of index entries required
     to fully index the document and publication system constraints.].
     protein gene sequence human cancer diagnosis therapy; cDNA protein
ST
     sequence human cancer diagnosis therapy
IT
     Gene, animal
     Gene, animal
      RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
      (Biological study)
         (, malignant tissue-specific expresssion and chromosome mapping of;
         human nucleic acids and corresponding proteins useful in the detection
         and treatment of various cancers)
     Gene, animal
IT
      RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
      (Biological study)
         (109P1D4, malignant tissue-specific expresssion and chromosome mapping
         of; human nucleic acids and corresponding proteins useful in the
         detection and treatment of various cancers)
      Gene, animal
IT
      RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
      (Biological study)
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(151P1C7A, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (151P4E11, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
TT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (154P2A8, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (156P1D4, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT
     Gene, animal
    RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (156P5C12, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
TΤ
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (159P2B5, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (161P2B7A, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
тт
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (179P3G7, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
     Gene, animal
IT
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (184P3C10B, malignant tissue-specific expresssion and chromosome
        mapping of; human nucleic acids and corresponding proteins useful in
        the detection and treatment of various cancers)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (184P3G10, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (185P2C9, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
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(185P3C2, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT
    Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (186P1H9, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
TT
    Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (187P3F2, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT.
    Gene. animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (192P2G7, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (74P3B3, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
TT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (83P4B8, malignant tissue-specific expresssion and chromosome mapping
        of; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
     Histocompatibility antigens
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-A1, epitopes; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
     Histocompatibility antigens.
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-A2, epitopes; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
     Histocompatibility antigens
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-A24, epitopes; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
     Histocompatibility antigens
TT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-A3, epitopes; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-B7, epitopes; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
     Histocompatibility antigens
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-DR3, epitopes; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
     PCR (polymerase chain reaction)
IT
        (RT-PCR (reverse transcription-PCR), diagnostic assay; human nucleic
        acids and corresponding proteins useful in the detection and treatment
        of various cancers)
TT
         (agents, conjugates with antibodies; human nucleic acids and
        corresponding proteins useful in the detection and treatment of various
        cancers)
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IT

Hybridoma

(antibody production by; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) TΤ Diagnosis Diagnosis (cancer; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) Intestine, neoplasm IT (colon; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) IT Cytotoxic agents (conjugates with antibodies; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) Antibodies and Immunoglobulins IT RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (conjugates, with cytotoxic agents or diagnostic reagents; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) IT T cell (lymphocyte) (cytotoxic, specific for tumor-restricted proteins; human nucleic acids and corresponding proteins useful in the detection and treatment of IT Immunoassay (diagnostic; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) Antibodies and Immunoglobulins IT RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (fragments; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) Antibodies and Immunoglobulins RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (fusion products; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) IT Genetic methods (gene discovery; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) IT T cell (lymphocyte) (helper cell, specific for tumor-restricted proteins; human nucleic acids and corresponding proteins useful in the detection and treatment of various cancers) IT Antitumor agents Bladder, neoplasm Epitopes Gene therapy Genetic mapping Human Kidney, neoplasm Kidney, neoplasm Lung, neoplasm Mammary gland, neoplasm Ovary, neoplasm Pancreas, neoplasm Population genetics Prostate gland, neoplasm Protein sequences Tumor markers

and treatment of various cancers) Tumor antigens

cDNA sequences

Tumor antigens RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)

(human nucleic acids and corresponding proteins useful in the detection

```
(human nucleic acids and corresponding proteins useful in the detection
        and treatment of various cancers)
IT
    Primers (nucleic acid)
    RL: ARG (Analytical reagent use); DGN (Diagnostic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
        (human nucleic acids and corresponding proteins useful in the detection
        and treatment of various cancers)
IT
    Antibodies and Immunoglobulins
    RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
    use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (human nucleic acids and corresponding proteins useful in the detection
        and treatment of various cancers)
IT
    Chromosome
        (human, gene mapping on; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
    Antibodies and Immunoglobulins
ΙT
    RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
    use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (humanized; human nucleic acids and corresponding proteins useful in
        the detection and treatment of various cancers)
    Antibodies and Immunoglobulins
IT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
    use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (monoclonal; human nucleic acids and corresponding proteins useful in
        the detection and treatment of various cancers)
IT
     Secondary structure
        (of tumor-restricted proteins; human nucleic acids and corresponding
        proteins useful in the detection and treatment of various cancers)
IT
     Plasmid vectors
        (polyepitopic; human nucleic acids and corresponding proteins useful in
        the detection and treatment of various cancers)
    Antibodies and Immunoglobulins
TT
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (single chain; human nucleic acids and corresponding proteins useful in
        the detection and treatment of various cancers)
IT
     Ribozymes
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (specific for tumor-restricted protein mRNA; human nucleic acids and
        corresponding proteins useful in the detection and treatment of various
        cancers)
IT
     B cell (lymphocyte)
     T cell (lymphocyte)
        (specific for tumor-restricted proteins; human nucleic acids and
        corresponding proteins useful in the detection and treatment of various
        cancers)
IT
     Animal
       (transgenic; human nucleic acids and corresponding proteins useful in
        the detection and treatment of various cancers)
IT
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 109P1D4; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
TT
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 151P1C7A; human nucleic acids and corresponding
        proteins useful in the detection and treatment of various cancers)
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
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(tumor-associated, 151P4E11; human nucleic acids and corresponding

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proteins useful in the detection and treatment of various cancers)
IT
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 154P2A8; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
TT
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 156P1D4; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
TТ
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 156P5C12; human nucleic acids and corresponding
        proteins useful in the detection and treatment of various cancers)
ΤТ
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses) (tumor-associated, 159P2B5; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
IT
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 161P2B7A; human nucleic acids and corresponding
        proteins useful in the detection and treatment of various cancers)
IT
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses) (tumor-associated, 179P3G7; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
IT
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 184P3C10B; human nucleic acids and corresponding
        proteins useful in the detection and treatment of various cancers)
IT
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 185P2C9; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
IT
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 185P3C2; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
IT
     Proteins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
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     BIOL (Biological study); USES (Uses)
        (tumor-associated, 187P3F2; human nucleic acids and corresponding proteins
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useful in the detection and treatment of various cancers)
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     BIOL (Biological study); USES (Uses)
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     BIOL (Biological study); USES (Uses)
        (tumor-associated, 192P2G7; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
IT
     Proteins
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     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
        (tumor-associated, 74P3B3; human nucleic acids and corresponding proteins
        useful in the detection and treatment of various cancers)
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     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
     BIOL (Biological study); USES (Uses)
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        useful in the detection and treatment of various cancers)
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     Vaccines
        (tumor; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
     Antitumor agents
IT
        (vaccines; human nucleic acids and corresponding proteins useful in the
        detection and treatment of various cancers)
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    BIOL (Biological study); USES (Uses)
        (peptide epitope; human nucleic acids and corresponding proteins useful
        in the detection and treatment of various cancers)
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     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
     use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
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BIOL (Biological study); USES (Uses)
        (peptide epitope; human nucleic acids and corresponding proteins useful
        in the detection and treatment of various cancers)
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     BIOL (Biological study); USES (Uses)
        (peptide epitope; human nucleic acids and corresponding proteins useful
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use); PRP (Properties); THU (Therapeutic use); ANST (Analytical study);
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     BIOL (Biological study); USES (Uses)
        (peptide epitope; human nucleic acids and corresponding proteins useful
        in the detection and treatment of various cancers)
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RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
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IT

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        (peptide epitope; human nucleic acids and corresponding proteins useful
        in the detection and treatment of various cancers)
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     L-Leucine, L-leucyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-threonyl-L-
CN
     leucyl-L-valyl-L-alanyl-L-leucyl-L-leucyl-L-seryl-L-leucylglycyl- (9CI)
     (CA INDEX NAME)
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Absolute stereochemistry.

PAGE 1-B

RN 473627-28-0 HCAPLUS

CN L-Serine, L-phenylalanyl-L-prolyl-L-prolyl-L-leucyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-threonyl-L-leucyl-L-valyl-L-alanyl-L-leucyl-L-leucyl-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

PAGE 2-B

RN 473627-44-0 HCAPLUS

CN L-Alanine, L-seryl-L-valyl-L-histidyl-L-phenylalanyl-L-prolyl-L-prolyl-L-leucyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-threonyl-L-leucyl-L-valyl-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B



PAGE 2-A

PAGE 3-A

RN 473628-58-9 HCAPLUS

CN L-Leucine, L-seryl-L-lysyl-L-seryl-L-valyl-L-histidyl-L-phenylalanyl-L-prolyl-L-prolyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-threonyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

Search done by Noble Jarrell

PAGE 1-B

PAGE 2-A

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L24 ANSWER 13 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
     2002:676325 HCAPLUS
     137:231363
DN
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     Entered STN: 08 Sep 2002
     Computerized methods for identifying T cell epitope and use for preparing
ΤI
     protein or antibody therapeutic agent with reduced immunogenicity
     Carr, Francis J.; Carter, Graham; Jones, Tim; Williams, Stephen; Hamilton,
IN
PA
     Merck Patent G.m.b.H., Germany
so
     PCT Int. Appl., 85 pp.
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
     ICM G06F019-00
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     15-3 (Immunochemistry)
     Section cross-reference(s): 1, 3, 63
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                 ECLA
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                        C07K016/30K; C07K016/46B2; C07K016/46B2B; G06F019/00C2
     Computerized method, with Bohm scoring function modified to include 12-6
AB
     van der Waal's ligand protein energy repulsive term and ligand
     conformational energy term based on MHC class II mol., is used for
     identification of T-cell epitopes, that give rise to an immune reaction in
     a living host. By means of this novel method biol. compds. can be
     generated which have a no or at least a reduced immunogenicity when
     exposed to the immune system of a given species and compared with the
     relevant non-modified entity. Thus the invention relates also to novel
     biol. mols., especially proteins and antibodies, obtained by the method
     according to the invention.
     computerized method T cell epitope protein antibody immunogenicity redn
st
IT
     Glycoproteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (40 kDa; computerized methods for identifying T cell epitope and use
        for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
IT
     Ricins
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RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (A, T cell epitope-modified; computerized methods for identifying T
        cell epitope and use for preparing protein or antibody therapeutic agent
        with reduced immunogenicity)
IT
    Apolipoproteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (A-I; computerized methods for identifying T cell epitope and use for
        preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
TΤ
    Antigens
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (A33; computerized methods for identifying T cell epitope and use for
        preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
TТ
     Proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (Acrp30; T cell epitope-modified; computerized methods for identifying
        T cell epitope and use for preparing protein or antibody therapeutic agent
        with reduced immunogenicity)
IT
     Antigens
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CD52; computerized methods for identifying T cell epitope and use for
        preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
TT
     Proteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (GLP-1; computerized methods for identifying T cell epitope and use for
        preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     Antibodies and Immunoglobulins
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (IgG1, T cell epitope-modified; computerized methods for identifying T
        cell epitope and use for preparing protein or antibody therapeutic agent
        with reduced immunogenicity)
     Histocompatibility antigens
TТ
     RL: ARU (Analytical role, unclassified); BSU (Biological study,
     unclassified); ANST (Analytical study); BIOL (Biological study)
        (MHC (major histocompatibility complex), binding epitope; computerized
        methods for identifying T cell epitope and use for preparing protein or
        antibody therapeutic agent with reduced immunogenicity)
ΙT
     Histocompatibility antigens
     RL: ARU (Analytical role, unclassified); BSU (Biological study,
     unclassified); ANST (Analytical study); BIOL (Biological study)
        (MHC (major histocompatibility complex), class II; computerized methods
        for identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
IT
     Biochemical compounds
        (T cell epitope-modified; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
     Antibodies and Immunoglobulins
IT
     Fusion proteins (chimeric proteins)
     Interleukin 1 receptor antagonist
     Proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (T cell epitope-modified; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
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TT
     Epitopes
        (T cell; computerized methods for identifying T cell epitope and use
        for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     Structure-activity relationship
IT
        (antibody-binding; computerized methods for identifying T cell epitope
        and use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     Structure-activity relationship
TT
        (antigen-binding; computerized methods for identifying T cell epitope
        and use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
TΤ
     Antigens
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (autoantigens, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
     Transplant and Transplantation
IT
        (bone marrow; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
ΙT
     Neurotrophic factors
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (brain-derived, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
IT
     Cytokines
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (cardiotrophin-like; T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
     Neurotrophic factor receptors
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (ciliary, T cell epitope-modified; computerized methods for identifying
        T cell epitope and use for preparing protein or antibody therapeutic agent
        with reduced immunogenicity)
TТ
     Anemia (disease)
     Antitumor agents
     Autoimmune disease
     Bacillus amyloliquefaciens
     Bacillus lentus
     Bacillus licheniformis
     Bacillus subtilis
     Blood vessel, disease
     Burn
     Cachexia
     Cardiopulmonary bypass
     Computer program
     Cosmetics
     Databases
     Detergents
     Diabetes mellitus
     Gaucher disease
     Hematopoiesis
     Human
     Hypertension
     Immunomodulators
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Immunosuppressants

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Infection
    Inflammation
    Injury
    Intestine, neoplasm
    Lymphoma
    Lysosomal storage disease
    Mus
    Pregnancy
    Protein sequences
    Rheumatoid arthritis
    Simulation and Modeling, physicochemical
    Surgery
    Textiles
    Thrombosis
    Transplant and Transplantation
    Transplant rejection
    Van der Waals potential
    Wound healing
        (computerized methods for identifying T cell epitope and use for preparing
        protein or antibody therapeutic agent with reduced immunogenicity)
IT
    CD20 (antigen)
    Epidermal growth factor receptors
     Interleukin 2 receptors
    neu (receptor)
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (computerized methods for identifying T cell epitope and use for preparing
        protein or antibody therapeutic agent with reduced immunogenicity)
ТT
     Ligands
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (conformational energy; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
IT
     Nerve, disease
        (degeneration; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
    Metabolism, animal
TΤ
        (disorder, cholesterol; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
IT
     Metabolism, animal
        (disorder, fatty acid; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
     Blood coagulation
IT
        (disorder; computerized methods for identifying T cell epitope and use
        for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
IT
     T cell (lymphocyte)
        (epitope; computerized methods for identifying T cell epitope and use
        for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
IT
     Kidney, disease
        (failure, chronic; computerized methods for identifying T cell epitope
        and use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     Interleukin 6
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (family; receptor cytokine-like factor 1; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
     Antibodies and Immunoglobulins
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
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PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (fragments, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
     Antibodies and Immunoglobulins
TТ
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (fusion products, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
     Transplant and Transplantation
IT
        (graft-vs.-host reaction; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
     Antibodies and Immunoglobulins
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (heavy chain, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
     Antibodies and Immunoglobulins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (humanized, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
TT
         (hypersensitivity; computerized methods for identifying T cell epitope
        and use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
IT
     Drug delivery systems
         (immunotoxins; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
ΙT
     Autoimmune disease
         (insulin-dependent diabetes mellitus; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
TT
     Diabetes mellitus
        (insulin-dependent; computerized methods for identifying T cell epitope
        and use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
IT
     Transplant and Transplantation
        (kidney, rejection; computerized methods for identifying T cell epitope
        and use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     Antibodies and Immunoglobulins
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (light chain, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
IT
     Fatty acids, biological studies
     RL: ADV (Adverse effect, including toxicity); BSU (Biological study,
     unclassified); BIOL (Biological study)
         (metabolic disease; computerized methods for identifying T cell epitope
        and use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
IT
     Antibodies and Immunoglobulins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
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(Preparation); USES (Uses)
        (monoclonal, T cell epitope-modified; computerized methods for
        identifying T cell epitope and use for preparing protein or antibody
        therapeutic agent with reduced immunogenicity)
IT
     Conformation
        (protein, ligand energy; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
IΤ
        (refractory; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     Mutagenesis
IT
        (site-directed, addition; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
     Mutagenesis
IT
        (site-directed, deletion; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
IT
     Mutagenesis
        (site-directed, substitution; computerized methods for identifying T
        cell epitope and use for preparing protein or antibody therapeutic agent
        with reduced immunogenicity)
IT
     Brain, disease
        (stroke; computerized methods for identifying T cell epitope and use
        for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
IT
     Eubacteria
     Fungi
     Mammalia
     Vertebrata
        (subtilisin; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
ΙT
     Kidney
        (transplant, rejection; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
TТ
        (transplant; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
ΙŢ
        (trauma; computerized methods for identifying T cell epitope and use
        for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     Tumor necrosis factor receptors
TΤ
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (type 1, T cell epitope-modified; computerized methods for identifying
        T cell epitope and use for preparing protein or antibody therapeutic agent
        with reduced immunogenicity)
     Tumor necrosis factor receptors
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (type 2; computerized methods for identifying T cell epitope and use
        for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
     148348-15-6P, Fibroblast growth factor 7
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
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(T cell epitope-modified; computerized methods for identifying T cell

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epitope and use for preparing protein or antibody therapeutic agent with
       reduced immunogenicity)
                                                           457106-95-5P
                    457106-94-4P, Leptin (human fragment)
TT
     457106-77-3P
     457106-96-6P
                    457106-97-7P, Erythropoietin (human fragment)
                                  457107-00-5P
                                                                 457107-02-7P
     457106-98-8P
                    457106-99-9P
                                                  457107-01-6P
     457107-03-8P
                    457107-04-9P
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                                                  457107-06-1P
                                                                 457107-07-2P
                    457107-09-4P
                                  457107-10-7P
                                                 457107-11-8DP, T-cell
     457107-08-3P
     epitope-modified derivs. 457107-12-9DP, T-cell epitope-modified derivs.
     457107-13-0DP, T-cell epitope-modified derivs.
                                                     457107-14-1DP, T-cell
                              457107-15-2DP, T-cell epitope-modified derivs.
     epitope-modified derivs.
     457107-16-3DP, T-cell epitope-modified derivs. 457107-17-4P
     457107-18-5P 457107-19-6P 457107-20-9P 457107-21-0P
                                                                457107-22-1DP,
    T-cell epitope-modified derivs. 457107-23-2DP, T-cell epitope-modified
              457107-24-3DP, T-cell epitope-modified derivs. 457107-25-4DP,
    derivs.
    T-cell epitope-modified derivs.
                                      457107-26-5P
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     457107-28-7P
                   457107-29-8P
    RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (amino acid sequence; computerized methods for identifying T cell
        epitope and use for preparing protein or antibody therapeutic agent with
        reduced immunogenicity)
IT
     65988-71-8, GD2
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (antibody to; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
TΤ
     80295-53-0, Complement C5
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (antibody to; computerized methods for identifying T cell epitope and
        use for preparing protein or antibody therapeutic agent with reduced
        immunogenicity)
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445388-46-5

445388-47-6

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     RL: ANT (Analyte); PRP (Properties); REM (Removal or disposal); ANST
     (Analytical study); PROC (Process)
        (computerized methods for identifying T cell epitope and use for preparing
        protein or antibody therapeutic agent with reduced immunogenicity)
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     RL: ANT (Analyte); PRP (Properties); REM (Removal or disposal); ANST
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(Analytical study); PROC (Process)
        (computerized methods for identifying T cell epitope and use for preparing
        protein or antibody therapeutic agent with reduced immunogenicity)
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     (Analytical study); PROC (Process)
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        protein or antibody therapeutic agent with reduced immunogenicity)
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RL: ANT (Analyte); PRP (Properties); REM (Removal or disposal); ANST
(Analytical study); PROC (Process)
   (computerized methods for identifying T cell epitope and use for preparing
   protein or antibody therapeutic agent with reduced immunogenicity)
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RL: ANT (Analyte); PRP (Properties); REM (Removal or disposal); ANST
(Analytical study); PROC (Process)
   (computerized methods for identifying T cell epitope and use for preparing
   protein or antibody therapeutic agent with reduced immunogenicity)
9014-01-1DP, Subtilisin, T cell epitope-modified derivs.
                                                            9024-58-2DP
Glutamic acid decarboxylase, isoforms and T cell epitope-modified derivs.
11096-26-7DP, Erythropoietin, T cell epitope-modified derivs.
37228-64-1DP, β-Glucocerebrosidase, T cell epitope-modified derivs.
60202-16-6P, Protein C
                        83869-56-1DP, GM-CSF, T cell epitope-modified
          143011-72-7DP, G-CSF, T cell epitope-modified derivs.
169494-85-3DP, Leptin, T cell epitope-modified derivs.
RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
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TT

IT

PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (computerized methods for identifying T cell epitope and use for preparing protein or antibody therapeutic agent with reduced immunogenicity)
126775-97-1D, Campath, T cell epitope-modified derivs. 152923-56-3D,
Zenapax, T cell epitope-modified derivs. 174722-31-7D, Rituxan, T cell epitope-modified derivs. 180288-69-1D, Herceptin, T cell epitope-modified derivs. 185243-69-0D, Enbrel, T cell epitope-modified derivs. 192391-48-3D, Bexxar, T cell epitope-modified derivs.
RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL

(Biological study); USES (Uses)
 (computerized methods for identifying T cell epitope and use for preparing protein or antibody therapeutic agent with reduced immunogenicity)
9002-68-0DP, Follicle stimulating hormone, T cell epitope-modified derivs.
RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);

PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(human: computerized methods for identifying T cell epitope and use for

(human; computerized methods for identifying T cell epitope and use for preparing protein or antibody therapeutic agent with reduced immunogenicity)

IT 57-88-5, Cholesterol, biological studies
RL: ADV (Adverse effect, including toxicity); BSU (Biological study, unclassified); BIOL (Biological study)

(metabolic disease associated with; computerized methods for identifying T cell epitope and use for preparing protein or antibody therapeutic agent with reduced immunogenicity)

IT 457873-85-7 457874-17-8 457874-75-8
RL: ANT (Analyte); PRP (Properties); REM (Removal or disposal); ANST (Analytical study); PROC (Process)

(computerized methods for identifying T cell epitope and use for preparing protein or antibody therapeutic agent with reduced immunogenicity)

RN 457873-85-7 HCAPLUS
CN L-Aspartic acid, L-histidyl-L-threonyl-L-valylglycyl-L-prolyl-L-histidyl-Lseryl-L-cysteinyl-L-histidyl-L-isoleucyl-L-prolyl-L-lysyl- (9CI) (CA
INDEX NAME)

Absolute stereochemistry.

IT

PAGE 1-A

PAGE 1-B

RN 457874-17-8 HCAPLUS

CN L-Phenylalanine, L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-histidyl-L-isoleucyl-L-prolyl-L-lysyl-L- α -aspartyl-L-leucyl-L-alanyl-L-leucyl-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

$$\begin{array}{c} & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\$$

PAGE 1-B

RN 457874-75-8 HCAPLUS

CN L-Isoleucine, L-α-glutamyl-L-α-glutamyl-L-tyrosyl-L-histidyl-L-threonyl-L-valylglycyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-histidyl-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

- L24 ANSWER 14 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
- AN 2002:631705 HCAPLUS
- DN 138:297158
- ED Entered STN: 22 Aug 2002
- TI Suppression of Tumor Recurrence and Metastasis by a Combination of the PHSCN Sequence and the Antiangiogenic Compound Tetrathiomolybdate in Prostate Carcinoma
- AU van Golen, Kenneth L.; Bao, Liwei; Brewer, George J.; Pienta, Kenneth J.; Kamradt, Jeffrey M.; Livant, Donna L.; Merajver, Sofia D.
- CS Division of Hematology and Oncology, Department of Internal Medicine, University of Michigan Comprehensive Cancer Center, Ann Arbor, MI,

Garcia 10/722843 48109-0948, USA Neoplasia (New York, NY, United States) (2002), 4(5), 373-379 SO CODEN: NEOPFL; ISSN: 1522-8002 Nature Publishing Group PB Journal DT English LА CC 1-6 (Pharmacology) Plasma fibronectin-mediated invasion of human DU145 prostate cancer cell AB line was efficaciously inhibited in a rat tumor model by treatment with Ac-PHSCN-NH2 peptide. Invasion of DU145 cells was stimulated by the PHSRN sequence of plasma fibronectin. However, PHSCN acts as a competitive inhibitor of PHSRN-mediated invasion. In the current study, we determined whether PHSCN could inhibit the recurrence and metastasis of DU145 tumors after excision of the primary tumor in an athymic nude mouse model. demonstrated that mice treated thrice weekly with i.v. Ac-PHSCN-NH2 peptide survived tumor-free for more than 30 wk post-primary tumor excision, whereas their untreated counterparts succumbed to recurrence and/or metastatic disease in significantly less time. Because of the universal requirement for angiogenesis in solid tumor growth, we tested the efficacy of copper deficiency induced by tetrathiomolybdate (TM) to retard tumor growth in the Dunning prostate cancer model. Significant reduction in size of the primary tumor was observed in mice rendered copper deficient. We sought to reduce tumor growth at the primary and metastatic sites by combining the anti-invasion Ac-PHSCN-NH2 peptide with TM. Improved survival, fewer metastatic lesions, and excellent tolerability were observed with the combination therapy. PHSCN peptide antiangiogenic tetrathiomolybdate prostate carcinoma STrecurrence metastasis Prostate gland, neoplasm IT (carcinoma, metastasis; suppression of tumor recurrence and metastasis by a combination of PHSCN sequence and the antiangiogenic compound tetrathiomolybdate in prostate carcinoma) Prostate gland, neoplasm (carcinoma; suppression of tumor recurrence and metastasis by a combination of PHSCN sequence and the antiangiogenic compound tetrathiomolybdate in prostate carcinoma) Carcinoma IT (prostatic, metastasis; suppression of tumor recurrence and metastasis by a combination of PHSCN sequence and the antiangiogenic compound tetrathiomolybdate in prostate carcinoma) IT Carcinoma (prostatic; suppression of tumor recurrence and metastasis by a combination of PHSCN sequence and the antiangiogenic compound tetrathiomolybdate in prostate carcinoma) IT Angiogenesis inhibitors Antitumor agents Drug interactions Human (suppression of tumor recurrence and metastasis by a combination of PHSCN sequence and the antiangiogenic compound tetrathiomolybdate in prostate carcinoma) TT 7440-50-8, Copper, biological studies

RL: BSU (Biological study, unclassified); BIOL (Biological study) (deficiency; efficacy of copper deficiency induced by tetrathiomolybdate (TM) to retard tumor growth in the Dunning prostate cancer model)

IT 16330-92-0 262438-43-7

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(suppression of tumor recurrence and metastasis by a combination of PHSCN sequence and the antiangiogenic compound tetrathiomolybdate in prostate carcinoma)

RE CNT 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD

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- (2) American Cancer Society; Cancer Facts and Figures 2002

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(3) Livant, D; Cancer Res 1995, V55, P5085 HCAPLUS
(4) Livant, D; Cancer Res 2000, V60, P309 HCAPLUS
(5) Mosher, D; Annu Rev Med 1984, V35, P561 HCAPLUS
(6) Nozue, M; Oncol Rep 2001, V8(6), P1247 MEDLINE
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(8) Rokhlin, O; Prostate 1995, V26, P205 HCAPLUS
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(11) Smith, D; Urol Clin North Am 1999, V26, P323 MEDLINE
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(13) Uchiyama, A; Br J Cancer 1999, V81(4), P721 MEDLINE
(14) van Golen, K; Clin Exp Metastasis 1996, V14, P95 HCAPLUS
(15) Webber, M; Clin Cancer Res 1995, V1, P1089 MEDLINE
(16) Witkowski, C; J Cancer Res Clin Oncol 1993, V119, P637 HCAPLUS
(17) Zheng, D; Cancer Res 1999, V59, P1655 HCAPLUS
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     L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl- (9CI)
CN
     (CA INDEX NAME)
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Absolute stereochemistry.

ANSWER 15 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

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2002:555761 HCAPLUS
ΑN
DN
     137:121939
ED
     Entered STN: 26 Jul 2002
     Compositions and methods for the use of fibronectin fragments in the
ΤI
     diagnosis of cancer
IN
     Livant, Donna
     The Regents of the University of Michigan, USA
PΔ
SO
     PCT Int. Appl., 77 pp.
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
     ICM G01N033-574
IC .
     ICS A61K049-00; A61K051-00
     9-16 (Biochemical Methods)
     Section cross-reference(s): 14
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                                               APPLICATION NO.
                                                                        DATE
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                                  DATE
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     WO 2002057786
                            A2
                                  20020725
                                               WO 2002-US1189
PΤ
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              GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
              LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
              PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
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                 CLASS PATENT FAMILY CLASSIFICATION CODES
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                         G01N033-574
                         A61K049-00; A61K051-00
                 ICS
                         A61K051/08; G01N033/50D4; G01N033/574V4
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 WO 2002057786
                 ECLA
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     MARPAT 137:121939
     The present invention concerns the detection tumors in vivo, the imaging
AΒ
     of tumors in vivo, and the imaging of cancerous tissue in pathol. samples.
     In particular the present invention incorporates the use of fibronectin
     fragments into these same detection and imaging methods.
     fibronectin cancer diagnosis imaging blood culture peptide radioisotope
ST
     fluorescent
IT
     Animal tissue
        (autopsy sample; compns. and methods for use of fibronectin fragments
        in diagnosis of cancer)
IT
     Radiography
        (autoradiography; compns. and methods for use of fibronectin fragments
        in diagnosis of cancer)
IT
     Animal tissue
     Blood analysis
     Blood plasma
     Blood serum
     Body fluid
     Diagnosis
     Epithelium
     Fibroblast
     Fluorescent substances
     Human
     Imaging
     Mammary gland, neoplasm
     Neoplasm
     Pathology
     Positron-emission tomography
     Prostate gland
     Rattus
     Sample preparation
     Scintigraphy
        (compns. and methods for use of fibronectin fragments in diagnosis of
        cancer)
     Peptides, uses
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     Radionuclides, uses
     RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical
     study); USES (Uses)
        (compns. and methods for use of fibronectin fragments in diagnosis of
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     Fibronectins
     RL: ANT (Analyte); BSU (Biological study, unclassified); DGN (Diagnostic
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        (degradation products; compns. and methods for use of fibronectin fragments
        in diagnosis of cancer)
IT
     Drug delivery systems
        (injections, i.v.; compns. and methods for use of fibronectin fragments
        in diagnosis of cancer)
IT
     Skin
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(keratinocyte; compns. and methods for use of fibronectin fragments in
        diagnosis of cancer)
ΙT
    Culture media
        (serum-free; compns. and methods for use of fibronectin fragments in
        diagnosis of cancer)
IT
        (tissue sample; compns. and methods for use of fibronectin fragments in
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        (compns. and methods for use of fibronectin fragments in diagnosis of
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     443305-23-5
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     study); BIOL (Biological study); USES (Uses)
        (compns. and methods for use of fibronectin fragments in diagnosis of
        cancer)
RN
     252230-05-0 HCAPLUS
     L-Asparagine, L-prolyl-L-histidyl-L-seryl-L-homocysteinyl- (9CI)
CN
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Absolute stereochemistry.

RN 262438-43-7 HCAPLUS

CN L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 443305-20-2 HCAPLUS

CN L-Asparagine, L-prolyl-L-histidyl-L-seryl-D-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 443305-23-5 HCAPLUS

CN L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-D-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 252229-85-9P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(compns. and methods for use of fibronectin fragments in diagnosis of cancer)

RN 252229-85-9 HCAPLUS

CN L-Asparagine, L-prolyl-L-histidyl-L-seryl-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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L24 ANSWER 16 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
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AN 2002:555624 HCAPLUS

DN 137:104821

ED Entered STN: 26 Jul 2002

TI Human nucleic acids encoding 50 human secreted proteins and their diagnostic and therapeutic uses

IN Moore, Paul A.; Ruben, Steven M.; Lafleur, David W.; Shi, Yanggu; Rosen, Craig A.; Olsen, Henrik; Ebner, Reinhard; Brewer, Laurie A.

PA Human Genome Sciences, Inc., USA

SO PCT Int. Appl., 785 pp.

CODEN: PIXXD2

DT Patent

LA English

IC ICM C12N

CC 3-3 (Biochemical Genetics)

Section cross-reference(s): 1, 6, 13

FAN.CNT 43 APPLICATION NO. DATE PATENT NO. KIND DATE ______ -----_____ _ _ _ _ ______ 20020117 <--20020725 WO 2002-US1109 ΡI WO 2002057420 A2 20021107 WO 2002057420 AЗ W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,

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WO 2002057420 ICM
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                      C07K014/47 ·
WO 2002057420 ECLA
    The present invention relates to 50 novel human secreted proteins and
     isolated nucleic acids containing the coding regions of the genes encoding
     such proteins. Also provided are vectors, host cells, antibodies, and
    recombinant methods for producing human secreted proteins. The invention
     further relates to diagnostic and therapeutic methods useful for
     diagnosing and treating diseases, disorders, and/or conditions related to
     these novel human secreted proteins.
ST
     secreted protein cDNA sequence human
TT
    Genetic mapping
        (chromosomal; human nucleic acids encoding 50 human secreted proteins
        and their diagnostic and therapeutic uses)
IT
    Disease, animal
        (diagnosis and treatment of secreted protein-associated; human nucleic
        acids encoding 50 human secreted proteins and their diagnostic and
        therapeutic uses)
IT
    Animal tissue
        (gene expression specificity in; human nucleic acids encoding 50 human
        secreted proteins and their diagnostic and therapeutic uses)
IT
     Drug screening
     Epitopes
     Human
    Molecular cloning
     Protein sequences
     cDNA sequences
        (human nucleic acids encoding 50 human secreted proteins and their
        diagnostic and therapeutic uses)
IT
     Antibodies and Immunoglobulins
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); THU (Therapeutic
     use); ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (human nucleic acids encoding 50 human secreted proteins and their
        diagnostic and therapeutic uses)
IT Gene, animal
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); PRP (Properties); BIOL (Biological study); USES (Uses)
        (human nucleic acids encoding 50 human secreted proteins and their
        diagnostic and therapeutic uses)
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     Chromosome
        (human, gene mapping on; human nucleic acids encoding 50 human secreted
        proteins and their diagnostic and therapeutic uses)
IT
        (mol.; human nucleic acids encoding 50 human secreted proteins and
        their diagnostic and therapeutic uses)
TТ
     Proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
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      Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
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    A single exon nucleic acid microarray comprising 12,614 single exon
AΒ
     nucleic acid probes for measuring gene expression in a sample derived from
     human lung cells is described. These unique exons are within longer probe
     sequences; sequencing confirms the exact chemical structure of each probe.
     Some amplicons have more than one exon, and some exons are contained in
     more than one amplicon. Expression, homol., and functional information
     are provided for the genome-derived single exon probes that are expressed
     significantly in human lung. Also described are 12,386 single exon
     nucleic acid probes and 12,011 proteins expressed in the lung and their
     use in methods for detecting gene expression. The genome-derived single
     exon nucleic acids comprise a novel type of nucleic acid microarray for
     verifying gene expression. In addition, methods are provided for identifying
     exons in a eukaryotic genome, and for assigning exons to a single gene.
     [This abstract record is one of nine records for this document necessitated
     by the large number of index entries required to fully index the document and
     publication system constraints.].
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     exon probe human lung
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     ANST (Analytical study); BIOL (Biological study); USES (Uses)
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Absolute stereochemistry.

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    Methods and apparatus for predicting, confirming and displaying functional
AB
     regions from genomic sequence data are used to identify 16,834 unique
     human genome-derived single exon probes useful for gene expression anal.,
     particularly gene expression anal. by microarray. Also presented are
     genome-derived single exon microarrays that include such probes, peptides
     encoded by the exons, and antibodies thereto. The human genome-derived
     single-exon probes are known to be expressed in one or more human tissues
     or cell types, particularly human brain, heart, liver, fetal
     liver, placenta, lung, bone marrow, BT474 and other human mammary
     epithelial cells, HeLa and other human cervical epithelial cells, and HBL
     100 and other human mammary epithelial cells. The invention provides a
     method of financing, selling and/or licensing genome-derived single-exon
     microarrays to customer desiring to measure gene expression, comprising:
     making available for computerized query or subscription service a database
     having a record corresponding to each genome-derived single exon
     microarray available for sale and/or license. [This abstract record is one
     of ten records for this document necessitated by the large number of index
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ST genome exon probe gene expression analysis; microarray exon probe gene expression analysis human; sequence gene exon probe human; business genome exon probe gene expression analysis

entries required to fully index the document and publication system

IT Animal cell line

(BT 474; human genome derived single exon nucleic acid probes useful for gene expression anal.)

IT Animal cell line

(HBL 100; human genome derived single exon nucleic acid probes useful for gene expression anal.)

IT Graph theory

(Mondrian representation of sequence and expression data; human genome derived single exon nucleic acid probes useful for gene expression anal.)

IT Liver

(adult and fetal; human genome derived single exon nucleic acid probes useful for gene expression anal.)

IT Industry

(bioinformatic; human genome derived single exon nucleic acid probes useful for gene expression anal.)

IT Uterus

(cervix, epithelium; human genome derived single exon nucleic acid probes useful for gene expression anal.)

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     Mammary gland
        (epithelium; human genome derived single exon nucleic acid probes
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IT
     Genetic element
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     Brain
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     Lung
     Molecular cloning
     Placenta
     Protein sequences
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IT
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IT
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TT
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                        C12Q001/68M6B; G06F019/00C1
     A single exon nucleic acid microarray comprising 5205 single exon nucleic
AΒ
     acid probes for measuring gene expression in a sample derived from human
     BT 474 cells is described. These unique exons are within longer probe
     sequences; sequencing confirms the exact chemical structure of each probe.
     Some amplicons have more than one exon, and some exons are contained in
     more than one amplicon. Expression, homol., and functional information
     are provided for the genome-derived single exon probes that are expressed
     significantly in human BT 474 cells. Also described are 5112 single exon
     nucleic acid probes and 5121 proteins expressed in the BT 474 cells and
     their use in methods for detecting gene expression. The genome-derived
     single exon nucleic acids comprise a novel type of nucleic acid microarray
     for verifying gene expression. In addition, methods are provided for
     identifying exons in a eukaryotic genome, and for assigning exons to a
     single gene. [This abstract record is one of three records for this
     document necessitated by the large number of index entries required to fully
     index the document and publication system constraints.].
     genome exon probe microarray gene expression BT474 cell; sequence genome
ST
     exon probe human breast
IT
     Mammary gland
        (BT 474 cells derived from; human genome-derived single exon nucleic
        acid probes useful for anal. of gene expression in human breast and BT
        474 cells)
IT
     Animal cell line
        (BT 474; human genome-derived single exon nucleic acid probes useful
        for anal. of gene expression in human breast and BT 474 cells)
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     Genetic element
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     ANST (Analytical study); BIOL (Biological study); USES (Uses)
        (exon; human genome-derived single exon nucleic acid probes useful for
        anal. of gene expression in human breast and BT 474 cells)
IT
     DNA microarray technology
     DNA sequences
     Protein sequences
        (human genome-derived single exon nucleic acid probes useful for anal.
        of gene expression in human breast and BT 474 cells)
IT
     Gene. animal
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     Human
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(Biological study)
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IT 400778-72-5

RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(amino acid sequence; human genome-derived single exon nucleic acid probes useful for anal. of gene expression in human breast and BT 474 cells)

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Absolute stereochemistry.

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PAGE 1-E

ANSWER 20 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

2001:624615 HCAPLUS AN

DN 135:328135

Entered STN: 29 Aug 2001

ED Nucleic acids containing single nucleotide polymorphisms in the human ΤI

Search done by Noble Jarrell

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genome
IN
     Shimkets, Richard A.; Leach, Martin
PA
     Curagen Corp., USA
     PCT Int. Appl., 4144 pp.
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     CODEN: PIXXD2
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     C07H021-04; C07H021-02; C12Q001-68; C07K014-47; C07K016-18; G01N033-53;
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     The invention provides 7867 nucleic acids containing single-nucleotide
     polymorphisms (SNPs) identified for transcribed human sequences, as well
     as methods of using the nucleic acids. The polymorphisms are arranged in
     the order: 5696 nucleotide sequences for SNPs that are silent; 315
     nucleotide sequences for SNPs that lead to conservative amino acid
     changes; 729 nucleotide changes for SNPs that lead to nonconservative
     amino acid changes; and 1127 nucleotide sequences for SNPs that involve a
     gap. The polymorphisms may be detected using allele-specific
     oligonucleotides that hybridize to the polymorphic site, and have
     applications in forensic analyses and disease diagnosis. [This abstract
     record is the second of 2 records for this document necessitated by the
     large number of index entries required to fully index the document and
     publication system constraints.].
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Forensic analysis
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   (unclaimed sequence; nucleic acids containing single nucleotide
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unclassified); BUU (Biological use, unclassified); PRP (Properties); ANST
(Analytical study); BIOL (Biological study); OCCU (Occurrence); USES
(Uses)
   (polymorphic site sequence; nucleic acids containing single nucleotide
   polymorphisms in the human genome)
364379-05-5 HCAPLUS
L\text{-Isoleucine, $L$-$\alpha$-glutamyl-$L$-valyl-$L$-prolyl-$L$-histidyl-$L$-seryl-$L$-
cysteinyl-L-arginyl-L-arginyl-L-tyrosyl-L-arginyl-L-leucyl-L-alanyl-L-
threonyl- (9CI) (CA INDEX NAME)
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Absolute stereochemistry.

IT

IT

RN

CN

PAGE 1-A

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PAGE 1-B

- L24 ANSWER 21 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
- AN 2001:472525 HCAPLUS
- DN 135:66204
- ED Entered STN: 29 Jun 2001
- TI A reversible linkage technology for controlled conjugation
- IN Flinn, Nicholas; Johnson, Tony
- PA Acambis Research Ltd., UK; Smithkline Beecham Biologicals SA
- SO PCT Int. Appl., 48 pp.
 - CODEN: PIXXD2
- DT Patent
- LA English
- IC ICM A61K047-48
- CC 63-5 (Pharmaceuticals)
 - Section cross-reference(s): 15

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             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
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                         A61K047/48R2V; A61K047/48R2L
                ECLA
 WO 2001045745
     MARPAT 135:66204
     The present invention describes a linkage for use in the conjugation of
     compds. (e.g. peptides) to carrier vehicles (e.g. macromols., polymers,
     dendrimers, proteins etc.), producing constructs of biol. and immunol.
     relevance. The ability to link an Epitope (such as a peptide) to a
     carrier (such as a protein) in a controlled and specific manner is of a
     paramount importance in the development of a potent, pharmaceutically
     relevant, immunogenic hapten-carrier construct, such as a vaccine. The
     invention provides a reversible linkage technol. for controlled
     conjugation of compds. such as peptides or peptidic Epitopes to carriers
     such as proteins. It utilizes an aryl aldehyde moiety to introduce an
     aldehyde functionality on to a carrier mol. (e.g. on to the surface of a
     protein carrier mol.). It uses a 2-hydroxy-4-alkoxy linker based on an
     aryl aldehyde to provide protection in the conjugation of a peptide to a
     carrier, by virtue of imine formation.
     vaccine epitope reversible linker aldehyde peptide
ST
     Proteins, specific or class
TΤ
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
         (D; reversible linkage technol. for controlled conjugation of epitopes
        to drug carriers)
IT
     Immunoglobulins
     RL: ADV (Adverse effect, including toxicity); BSU (Biological study,
     unclassified); BIOL (Biological study)
         (E, -mediated disease; reversible linkage technol. for controlled
        conjugation of epitopes to drug carriers)
IT
     Immunostimulants
         (adjuvants; reversible linkage technol. for controlled conjugation of
        epitopes to drug carriers)
     Drug delivery systems
IT
         (carriers; reversible linkage technol. for controlled conjugation of
        epitopes to drug carriers)
ΙT
     Peptides, biological studies
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); THU
      (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
         (cyclic, epitopes; reversible linkage technol. for controlled
        conjugation of epitopes to drug carriers)
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
         (diphtheria; reversible linkage technol. for controlled conjugation of
        epitopes to drug carriers)
IT
     Proteins, general, biological studies
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RL: BPR (Biological process); BSU (Biological study, unclassified); PEP
     (Physical, engineering or chemical process); THU (Therapeutic use); BIOL
     (Biological study); PROC (Process); USES (Uses)
        (drug carriers; reversible linkage technol. for controlled conjugation
        of epitopes to drug carriers)
     Peptides, biological studies
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     RL: BPR (Biological process); BSU (Biological study, unclassified); PEP
     (Physical, engineering or chemical process); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (epitopes; reversible linkage technol. for controlled conjugation of
        epitopes to drug carriers)
IT
     Fissurella
        (hemocyanin of; reversible linkage technol. for controlled conjugation
        of epitopes to drug carriers)
IT
    Toxins
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
        (inactivated bacterial; reversible linkage technol. for controlled
        conjugation of epitopes to drug carriers)
     Drug delivery systems
TT
        (liposomes; reversible linkage technol. for controlled conjugation of
        epitopes to drug carriers)
IT
    Hemocyanins
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
        (of keyhole limpet; reversible linkage technol. for controlled
        conjugation of epitopes to drug carriers)
    Drug delivery systems
IT
     Epitopes
     Immunostimulants
     Molecular cloning
     Phage display
     Vaccines
        (reversible linkage technol. for controlled conjugation of epitopes to
        drug carriers)
IT
    Antibodies
     RL: BPR (Biological process); BSU (Biological study, unclassified); MFM
     (Metabolic formation); THU (Therapeutic use); BIOL (Biological study);
     FORM (Formation, nonpreparative); PROC (Process); USES (Uses)
        (reversible linkage technol. for controlled conjugation of epitopes to
        drug carriers)
IT
     Antigens
     Tuberculin
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
        (reversible linkage technol. for controlled conjugation of epitopes to
        drug carriers)
TТ
     Cattle
        (serum albumin of; reversible linkage technol. for controlled
        conjugation of epitopes to drug carriers)
     Albumins, biological studies
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     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
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        epitopes to drug carriers)
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     Toxins
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (tetanus; reversible linkage technol. for controlled conjugation of
        epitopes to drug carriers)
IT
     Bacteria (Eubacteria)
        (toxins of; reversible linkage technol. for controlled conjugation of
        epitopes to drug carriers)
IT
     Protein motifs
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        conjugation of epitopes to drug carriers)
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7429-90-5D, Aluminum, salts, biological studies 7440-70-2D, Calcium,
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     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); PEP (Physical, engineering or chemical process); THU
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IT 6066-82-6 206656-20-4

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IT 346460-13-7P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

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Absolute stereochemistry.

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     Cloning and cDNA and deduced amino acid sequences of 52 human secreted
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     Human Genome Sciences, Inc., USA; et al.
PA
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AB
     isolated nucleic acids containing the coding regions of the genes encoding
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     epitope sites are provided for the secreted proteins, as well as
     chromosomal mapping of some of the genes. Also provided are vectors, host
     cells, antibodies, and recombinant methods for producing human secreted
     proteins in bacterial, insect, and mammalian cells. The invention further
     relates to diagnostic and therapeutic methods useful for diagnosing and
     treating disorders related to these novel human secreted proteins.
     High-throughput screening assays are also provided for various putative
     activities of the secreted proteins.
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     Animal cell line
IT
        (CHO, recombinant host; cloning and cDNA and deduced amino acid
        sequences of 52 human secreted proteins)
     Animal cell line
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        (SF9, recombinant host; cloning and cDNA and deduced amino acid
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        (chromosomal mapping of secreted protein genes; cloning and cDNA and
        deduced amino acid sequences of 52 human secreted proteins)
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     Epitopes
     Gene therapy
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        (cloning and cDNA and deduced amino acid sequences of 52 human secreted
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(1) Agostino; Database GenBank (N_Geneseq 36); Accesion NO: V86545 1998
(2) Hillier; Database GenBank (EST); Accession NO: H08444 1995
(3) Matsubara; Database GenBank (N_Geneseq_36); Accession NO: T23968 1995
(4) NCI-CGAP; Database GenBank (EST); Accession NO:AI92802 1999
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Absolute stereochemistry.

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PAGE 1-B

PAGE 1-C

PAGE 1-E

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PAGE 2-D

Dh

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    A method for protecting a peptide from peptidase activity in vivo, the
     peptide being composed of between 2 and 50 amino acids and having a
    C-terminus and an N-terminus and a C-terminus amino acid and an N-terminus
    amino acid is described. In the first step of the method, the peptide is
    modified by attaching a reactive group to the C-terminus amino acid, to
     the N-terminus amino acid, or to an amino acid located between the
    N-terminus and the C-terminus, such that the modified peptide is capable
     of forming a covalent bond in vivo with a reactive functionality on a
    blood component. The solid phase peptide synthesis of a number of derivs.
    with 3-maleimidopropionic acid (3-MPA) is described. In the next step, a
     covalent bond is formed between the reactive group and a reactive
     functionality on a blood component to form a peptide-blood component
     conjugate, thereby protecting said peptide from peptidase activity. The
     final step of the method involves the analyzing of the stability of the
    peptide-blood component conjugate to assess the protection of the peptide
     from peptidase activity. Thus, the percentage of a K5 kringle peptide
     (Pro-Arg-Lys-Leu-Tyr-Asp-Lys-NH2) conjugated to human serum albumin via
    MPA remained relatively constant through a 24-h plasma assay in contrast to
     unmodified K5 which decreased to 9% of the original amount of K5 in only 4 h
     in plasma.
     therapeutic peptide conjugation blood albumin stability; peptidase
     stability therapeutic peptide conjugate blood; maleimido conjugation
     therapeutic peptide peptidase stability
     Proteins, specific or class
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN
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     PREP (Preparation); PROC (Process); USES (Uses)
        (CART (cocaine and amphetamine-regulated transcript); protection of
        endogenous therapeutic peptides from peptidase activity through
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     PREP (Preparation); PROC (Process); USES (Uses)
        (apoptosis-regulating; protection of endogenous therapeutic peptides
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IT
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IT
     Peptides, biological studies
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN
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     PREP (Preparation); PROC (Process); USES (Uses)
        (conjugates, therapeutic; protection of endogenous therapeutic peptides
        from peptidase activity through conjugation to blood components)
IT
     Fibrinogen degradation products
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN
     (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study);
     PREP (Preparation); PROC (Process); USES (Uses)
(fibrinopeptides; protection of endogenous therapeutic peptides from
        peptidase activity through conjugation to blood components)
     Fibronectins
     Laminins
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study);
     PREP (Preparation); PROC (Process); USES (Uses)
        (fragments; protection of endogenous therapeutic peptides from
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peptidase activity through conjugation to blood components)
IT
     Complement
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN
     (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study);
     PREP (Preparation); PROC (Process); USES (Uses)
        (inhibitors; protection of endogenous therapeutic peptides from
        peptidase activity through conjugation to blood components)
TΤ
     Functional groups
        (maleimido or succinimidyl; protection of endogenous therapeutic
        peptides from peptidase activity through conjugation to blood
TТ
     Solid phase synthesis
        (peptide; protection of endogenous therapeutic peptides from peptidase
        activity through conjugation to blood components)
IT
    Antimicrobial agents
     Antioxidants
     Signal transduction, biological
        (peptides; protection of endogenous therapeutic peptides from peptidase
        activity through conjugation to blood components)
    Cell adhesion molecules
TT
    Chemotactic factors
     G proteins (guanine nucleotide-binding proteins)
     Gastrointestinal hormones
     Hypothalamic hormones
     Interleukin receptors
     Interleukins
     Osteocalcins
     Pancreatic hormones
     Pituitary hormones
     Tachykinins
     Thyroid hormones
     Toxins
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN
     (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study);
     PREP (Preparation); PROC (Process); USES (Uses)
        (protection of endogenous therapeutic peptides from peptidase activity
        through conjugation to blood components)
     Albumins, biological studies
     RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological
     study); PREP (Preparation); USES (Uses)
        (serum, conjugates; protection of endogenous therapeutic peptides from
        peptidase activity through conjugation to blood components)
IT
     Amyloid
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN
     (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study);
     PREP (Preparation); PROC (Process); USES (Uses)
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        peptidase activity through conjugation to blood components)
IT
     124219-00-7
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        peptidase activity through conjugation to blood components)
IT
     169494-85-3P, Leptin
     RL: BPR (Biological process); BSU (Biological study, unclassified); SPN
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     PREP (Preparation); PROC (Process); USES (Uses)
        (fragments; protection of endogenous therapeutic peptides from
        peptidase activity through conjugation to blood components)
     9002-04-4P, Thrombin
TT
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     PREP (Preparation); PROC (Process); USES (Uses)
        (inhibitors; protection of endogenous therapeutic peptides from
        peptidase activity through conjugation to blood components)
IT
     9031-96-3, Peptidase
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
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study, unclassified); BIOL (Biological study)
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     50-56-6P, Oxytocin, biological studies 1393-25-5P, Secretin
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     Gastrin
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     57285-09-3P, Inhibin
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     80043-53-4P, Gastrin-releasing peptide 81858-94-8P, Peptide YY (swine)
     82785-45-3P, Neuropeptide Y 83652-28-2P, Calcitonin gene related peptide
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              106602-62-4P, Amylin
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     116110-78-2P, Insulin-like peptide 117148-67-1P, Pancreastatin
     119418-04-1P, Galanin 127830-04-0P, C-Type natriuretic peptide 137061-48-4P, Pituitary adenylate cyclase-activating polypeptide
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     6066-82-6, N-Hydroxysuccinimide
IT
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         (substrates and inhibitors; protection of endogenous therapeutic
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peptides from peptidase activity through conjugation to blood components) 11063-17-5; Gastric inhibitory polypeptide (swine major) 12279-41-3, IT α1-39-Corticotropin (human) 51257-86-4 52232-67-4 68563-24-6, 68893-82-3 Corticotropin-inhibiting peptide (human) 75718-92-2 Peptide F (cattle adrenal medulla) 77465-10-2, α1-39-Corticotropin 82707-63-9 83328-95-4, Dynorphin-32 (swine) 84069-10-3. 1-40-Somatoliberin (human pancreatic islet) 84136-38-9, 1-37-Somatoliberin (human pancreatic islet) 85568-24-7, 86472-71-1, Somatoliberin (rat hypothalamus) 1-44-Parathormone (human) 87713-86-8, Peptide B (cattle adrenal medulla) 87805-34-3, Glucagon-like peptide I (human) 88402-03-3, Defensin NP 3A (rabbit reduced) 89747-89-7, Transforming growth factor α (rat reduced) 89492-47-7 92879-83-9, α -Inhibin-31 (human seminal 90599-39-6 90880-43-6 98614-76-7 99120-49-7, 96351-01-8 96827-07-5 98226-75-6 plasma) 99658-10-3 100040-31-1, Glucagon-like peptide II (human) 99658-03-4 100915-92-2 101178-74-9 Gastric inhibitory polypeptide (human) 107761-42-2, Glycopeptide (human clone 9-110 amyloid A4 peptide moiety) 110616-04-1 · 111274-30-7, Pancreatic polypeptide (Canis familiaris) 112540-82-6 112199-06-1 112602-83-2, Somatoliberin 111366-38-2 116977-48-1, Defensin NP 4 (human reduced) 112938-42-8 (carp) 119977-20-7 118337-11-4, Echistatin α 1 (reduced) 118277-01-3 121181-17-7, Glucagon-like peptide 1 (Octodon degus) 120298-73-9 123337-90-6, Brain natriuretic peptide-45 (rat reduced) 122931-92-4 124147-28-0, Lymphokine CINC (rat subunit reduced) 123757-43-7 125199-49-7, Somatoliberin 124834-82-8 124834-83-9 124541-50-0 127609-05-6 128906-76-3, Decorsin 126466-47-5 126466-48-6 (mouse) 131438-79-4 134090-87-2 131438-74-9 (reduced) 129405-61-4 134710-25-1, Calciseptin (Dendroaspis polylepis polylepis 134500-80-4 136751-54-7, C-Type natriuretic peptide (human reduced) 135842-15-8 137467-72-2, C-Type clone λhCNP 53-amino acid isoform reduced) natriuretic peptide (swine clone λCNP6 53-amino acid isoform 142192-88-9 138634-59-0 139872-85-8 141983-93-9 144409-98-3 144409-99-4 144855-49-2 146103-03-9 142297-36-7 148846-98-4, Brain natriuretic peptide-45 (mouse reduced) 150679-91-7 150871-76-4 150680-28-7, Adrenomedullin (human clone pHAM-3) 151247-82-4, Margatoxin (reduced) 154101-11-8 155420-59-0 161246-56-6 161246-49-7 161246-57-7 155924-66-6 157938-23-3 161246-63-5 161246-62-4 161246-58-8 161246-60-2 161246-61-3 161246-69-1 161246-68-0 161246-72-6 161246-74-8 161246-67-9 161246-92-0 161246-93-1 161246-94-2 161246-75-9 161246-87-3 161247-04-7 161247-07-0 161247-08-1 161247-09-2 161247-06-9 161247-16-1 161247-12-7 161247-21-8 161247-23-0 161247-11-6 161247-24-1 161247-34-3 161247-35-4 161247-37-6 161247-48-9 161247-61-6 161247-58-1 161247-49-0 161247-50-3 161247-56-9 161818-07-1, Endothelin 3 (rat reduced) 164714-89-0, 161278-54-2 ω-Agatoxin TK 2 (reduced) 165168-50-3, Toxin ShK (Stoichactis helianthus potassium-channel reduced) 166090-74-0 166798-69-2 171543-83-2, Urocortin (Rattus norvegicus) 173010-28-1, Buforin I 175524-68-2, Glucagon-like peptide 1 (Amphiuma tridactylum) 174394-41-3 175779-77-8, Glucagon-like peptide 2 (Amphiuma tridactylum) 179733-43-8 183598-56-3 183681-18-7 184007-69-0 186322-91-8 179733-44-9 186673-16-5 186673-15-4 186673-13-2 186673-14-3 186673-12-1 186673-19-8 186673-20-1 186673-21-2 186673-18-7 186673-17-6 187111-91-7 187547-59-7 186811-41-6 187111-84-8 187111-86-0 188762-99-4 192588-09-3 192588-12-8 195262-56-7 203265-23-0 213779-10-3 208540-81-2 211496-36-5 203743-40-2 206010-80-2 227015-70-5 235787-61-8 256345-50-3 213779-21-6 221377-92-0 256345-97-8 256345-99-0 256345-94-5 256345-89-8 256345-87-6 256346-54-0 256346-89-1 256346-01-7 256346-36-8 256346-45-9 306277-48-5 303052-45-1 256346-91-5 260363-52-8 302905-40-4 309255-24-1 309255-30-9 309255-32-1 309255-20-7 308388-50-3 309255-44-5 309255-62-7 309255-64-9 309255-50-3 309255-60-5 309255-73-0) 309255-72-9 309255-69-4 309255-71-8 309255-68-3 309255-77-4 309255-78-5 309255-74-1 309255-75-2 309255-76-3 309256-41-5 309255-81-0 309255-91-2 309255-79-6 309255-80-9

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(swine)

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Absolute stereochemistry.

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L24 ANSWER 24 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN AN 2000:688253 HCAPLUS
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DN 133:248082

ED Entered STN: 29 Sep 2000

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Cloning and cDNA and deduced amino acid sequences of 49 human secreted
TI
     proteins
     Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George
IN
     Human Genome Sciences, Inc., USA
PΑ
     PCT Int. Appl., 431 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
     ICM C07H021-04
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     ICS C07K014-00; C07K016-00; C12N015-00; C12N015-63; C12N015-85;
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     3-3 (Biochemical Genetics)
     Section cross-reference(s): 6, 13, 63
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         AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
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                  ICS
                         C12N015-85; C12N015-86; C12Q001-68; G01N033-53
 WO 2000056753 ECLA
                         C07K014/47
     The present invention relates to 49 novel human secreted proteins and
AB
     isolated nucleic acids containing the coding regions of the genes encoding
     such proteins. Tissue distribution, sequence homologies, and preferred
     epitope sites are provided for the secreted proteins, as well as
     chromosomal mapping of some of the genes. Also provided are vectors, host
     cells, antibodies, and recombinant methods for producing human secreted
     proteins in bacterial, insect, and mammalian cells. The invention further
     relates to diagnostic and therapeutic methods useful for diagnosing and
     treating disorders related to these novel human secreted proteins.
     High-throughput screening assays are also provided for various putative
     activities of the secreted proteins.
     secretory protein cDNA sequence human
ST
     Animal cell line
IT
         (CHO, recombinant host; cloning and cDNA and deduced amino acid
         sequences of 49 human secreted proteins)
IT
     Genetic element
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
         (GAS (gamma activation site), Jaks-STAT pathway for high-throughput
         screening assays; cloning and cDNA and deduced amino acid sequences of
         49 human secreted proteins)
     Genetic element
TΤ
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
         (IRS (interferon-responsive sequence), Jaks-STAT pathway for
        high-throughput screening assays; cloning and cDNA and deduced amino
         acid sequences of 49 human secreted proteins)
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IT
     Signal transduction, biological
        (Jaks-STAT pathway for high-throughput screening assays; cloning and
        cDNA and deduced amino acid sequences of 49 human secreted proteins)
     Transcription factors
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     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (NF-κB (nuclear factor κB), high-throughput screening
        assays; cloning and cDNA and deduced amino acid sequences of 49 human
     secreted proteins)
Animal cell line
TT
        (SF9, recombinant host; cloning and cDNA and deduced amino acid
        sequences of 49 human secreted proteins)
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     (Biological study); PROC (Process)
        (STAT, Jaks-STAT pathway for high-throughput screening assays; cloning
        and cDNA and deduced amino acid sequences of 49 human secreted
        proteins)
     Genetic mapping
TT
        (chromosomal mapping of secreted protein genes; cloning and cDNA and
        deduced amino acid sequences of 49 human secreted proteins)
IT
     Drugs
     Epitopes
     Gene therapy
     Molecular cloning
        (cloning and cDNA and deduced amino acid sequences of 49 human secreted
        proteins)
IT
     Antibodies
     RL: ARG (Analytical reagent use); THU (Therapeutic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
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TΤ
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     BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES
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        (cloning and cDNA and deduced amino acid sequences of 49 human secreted
        proteins)
     Signal peptides
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     (Properties); BIOL (Biological study); OCCU (Occurrence)
        (cloning and cDNA and deduced amino acid sequences of 49 human secreted
        proteins)
IT
     Diagnosis
     Mutation
        (disease diagnosis by mutation detection; cloning and cDNA and deduced
        amino acid sequences of 49 human secreted proteins)
IT
     cDNA sequences
        (for 49 human secreted proteins)
IT
     Immunoglobulins
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        (fusion products; cloning and cDNA and deduced amino acid sequences of
        49 human secreted proteins)
     T cell (lymphocyte)
TТ
        (high-throughput screening assays for T-cell activity; cloning and cDNA
        and deduced amino acid sequences of 49 human secreted proteins)
IT
     Chromosome
        (human, chromosomal mapping of secreted protein genes; cloning and cDNA
        and deduced amino acid sequences of 49 human secreted proteins)
     Animal cell
IT
        (mammalian, recombinant host; cloning and cDNA and deduced amino acid
        sequences of 49 human secreted proteins)
ΙT
     Hematopoietic precursor cell
        (myeloid, high-throughput screening assays for myeloid activity;
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cloning and cDNA and deduced amino acid sequences of 49 human secreted

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proteins)
TT
    Nerve
        (neuron, high-throughput screening assays for neuronal activity;
        cloning and cDNA and deduced amino acid sequences of 49 human secreted
       proteins)
    Protein sequences
IT
        (of 49 human secreted proteins)
TT
    Biological transport
        (permeation, high-throughput screening assays; cloning and cDNA and
        deduced amino acid sequences of 49 human secreted proteins)
    Phosphorylation, biological
IT
        (protein, high-throughput screening assays; cloning and cDNA and
        deduced amino acid sequences of 49 human secreted proteins)
IT
    Escherichia coli
        (recombinant host; cloning and cDNA and deduced amino acid sequences of
        49 human secreted proteins)
    Proteins, specific or class
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     RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU
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(1) Isomura; Database GenEmbl, section 2/19 2000, AP000066
(2) Marra; Database EST 1998, AI042727
(3) Strausberg; Database EST 1998, AI285709
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Absolute stereochemistry.

L24 ANSWER 25 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2000:666747 HCAPLUS

DN 133:233609

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ED
     Entered STN: 22 Sep 2000
     Cloning and cDNA and deduced amino acid sequences of 50 human secreted
ΤI
     proteins
IN
     Rosen, Craig A.; Ruben, Steven M.; Komatsoulis, George
     Human Genome Sciences, Inc., USA
PA
     PCT Int. Appl., 428 pp.
SO
     CODEN: PIXXD2
DT
     Patent
     English
LΑ
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     ICM C07H021-04
     ICS C07K014-00; C07K016-00; C12N015-00; C12N015-63; C12N015-85;
          C12N015-86; C12Q001-68; G01N033-53
     3-3 (Biochemical Genetics)
     Section cross-reference(s): 6, 13, 63
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                        C07K014-00; C07K016-00; C12N015-00; C12N015-63;
                 ICS
                         C12N015-85; C12N015-86; C12Q001-68; G01N033-53
 WO 2000055175 ECLA
                        C07K014/47
     The present invention relates to 50 novel human secreted proteins and
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     isolated nucleic acids containing the coding regions of the genes encoding
     such proteins. Tissue distribution, sequence homologies, and preferred
     epitope sites are provided for the secreted proteins, as well as
     chromosomal mapping of some of the genes. Also provided are vectors, host
     cells, antibodies, and recombinant methods for producing human secreted
     proteins in bacterial, insect, and mammalian cells. The invention further
     relates to diagnostic and therapeutic methods useful for diagnosing and
     treating disorders related to these novel human secreted proteins.
     High-throughput screening assays are also provided for various putative
     activities of the secreted proteins.
     secretory protein cDNA sequence human
ST
     Animal cell line
IT
         (CHO, recombinant host; cloning and cDNA and deduced amino acid
        sequences of 50 human secreted proteins)
ΙT
     Genetic element
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
         (GAS (gamma activation site), Jaks-STAT pathway for high-throughput
        screening assays; cloning and cDNA and deduced amino acid sequences of
        50 human secreted proteins)
     Genetic element
IT
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        (IRS (interferon-responsive sequence), Jaks-STAT pathway for
        high-throughput screening assays; cloning and cDNA and deduced amino
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acid sequences of 50 human secreted proteins)
     Signal transduction, biological
        (Jaks-STAT pathway for high-throughput screening assays; cloning and
        cDNA and deduced amino acid sequences of 50 human secreted proteins)
    Transcription factors
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    RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (NF-κB (nuclear factor κB), high-throughput screening
        assays; cloning and cDNA and deduced amino acid sequences of 50 human
        secreted proteins)
    Animal cell line
IT
        (SF9, recombinant host; cloning and cDNA and deduced amino acid
        sequences of 50 human secreted proteins)
IT
    Transcription factors
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (STAT, Jaks-STAT pathway for high-throughput screening assays; cloning
        and cDNA and deduced amino acid sequences of 50 human secreted
        proteins)
TT
     Genetic mapping
        (chromosomal mapping of secreted protein genes; cloning and cDNA and
        deduced amino acid sequences of 50 human secreted proteins)
LT
    Drugs
     Epitopes
     Gene therapy
    Molecular cloning
        (cloning and cDNA and deduced amino acid sequences of 50 human secreted
        proteins)
    Antibodies
TT
     RL: ARG (Analytical reagent use); THU (Therapeutic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
        (cloning and cDNA and deduced amino acid sequences of 50 human secreted
        proteins)
TΤ
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     BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES
        (cloning and cDNA and deduced amino acid sequences of 50 human secreted
        proteins)
TT
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     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); PRP
     (Properties); BIOL (Biological study); OCCU (Occurrence)
        (cloning and cDNA and deduced amino acid sequences of 50 human secreted
        proteins)
IT
     Diagnosis
     Mutation
        (disease diagnosis by mutation detection; cloning and cDNA and deduced
        amino acid sequences of 50 human secreted proteins)
IT
     cDNA sequences
        (for 50 human secreted proteins)
IT
     Immunoglobulins
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (fusion products; cloning and cDNA and deduced amino acid sequences of
        50 human secreted proteins)
IT
     T cell (lymphocyte)
        (high-throughput screening assays for T-cell activity; cloning and cDNA
        and deduced amino acid sequences of 50 human secreted proteins)
IT
     Chromosome
        (human, chromosomal mapping of secreted protein genes; cloning and cDNA
        and deduced amino acid sequences of 50 human secreted proteins)
IT
     Animal cell
        (mammalian, recombinant host; cloning and cDNA and deduced amino acid
        sequences of 50 human secreted proteins)
IT
     Hematopoietic precursor cell
        (myeloid, high-throughput screening assays for myeloid activity;
```

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cloning and cDNA and deduced amino acid sequences of 50 human secreted
        proteins)
IT
     Nerve
        (neuron, high-throughput screening assays for neuronal activity;
        cloning and cDNA and deduced amino acid sequences of 50 human secreted
        proteins)
     Protein sequences
IT
        (of 50 human secreted proteins)
IT
     Biological transport
        (permeation, high-throughput screening assays; cloning and cDNA and
        deduced amino acid sequences of 50 human secreted proteins)
     Phosphorylation, biological
IT
        (protein, high-throughput screening assays; cloning and cDNA and
        deduced amino acid sequences of 50 human secreted proteins)
IT
     Escherichia coli
        (recombinant host; cloning and cDNA and deduced amino acid sequences of
        50 human secreted proteins)
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IT
     RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU
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     BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES
        (secretory; cloning and cDNA and deduced amino acid sequences of 50
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(1) Anon; Database GENEMBL
(2) Anon; Database GENEMBL
(3) Blum; J Biol Chem 1996, V271(29), P17183 HCAPLUS
(4) Blum; J Biol Chem 1996, V271(29), P17183 HCAPLUS
(5) Holthuis; Biochem J 1995, V312, P205 HCAPLUS
(6) Rojo; J Cell Biol 1997, V139(5), P1119 HCAPLUS
(7) Sanseau; Direct Submission 1996
(8) Sherrington; Nature 1995, V375(6534), P754 HCAPLUS
(9) Sohn; J Cell Biol 1996, V135(5), P1239 HCAPLUS
(10) Trower; Proc Natl Acad Sci U S A 1996, V93, P1366 HCAPLUS
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Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

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PAGE 1-C

PAGE 2-C

L24 ANSWER 26 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2000:608780 HCAPLUS

DN 133:206771

ED Entered STN: 01 Sep 2000

TI Epitopes or mimotopes derived from the CE2 domain of IgE, antagonists thereof, and their therapeutic uses

IN Dyson, Michael; Friede, Martin; Greenwood, Judith; Hewitt, Ellen; Lamont, Alan; Mason, Sean; Randall, Roger; Turnell, William Gordon; Van Mechelen, Marcelle Paulette; Vinals y De Bassols, Carlota

PA Smithkline Beecham Biologicals SA, Belg.; Peptide Therapeutics Limited

SO PCT Int. Appl., 129 pp.

CODEN: PIXXD2

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     ICM C07K016-00
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           G01N033-68; A61P037-08
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                          A61K047/48R2V; A61K047/48R2H; C07K016/00; C07K016/42M10
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                          4C085/EE06; 4C085/FF24; 4H045/AA10; 4H045/AA11;
                          4H045/AA20; 4H045/BA10; 4H045/BA13; 4H045/BA14;
                          4H045/BA15; 4H045/BA16; 4H045/BA17; 4H045/BA41;
                          4H045/CA40; 4H045/DA76; 4H045/DA86; 4H045/EA28;
                          4H045/EA50; 4H045/FA34; 4H045/FA44; 4H045/FA51;
                          4H045/FA72; 4H045/FA74; 4H045/GA22
     The present invention relates to the provision of novel medicaments for
AB
     the treatment, prevention or amelioration of allergic disease. In
     particular, the novel medicaments are isolated peptides incorporating
     epitopes or mimotopes of surface exposed regions of the C<e2 domain of
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IGE. The inventors have found that these novel regions may be the target
     for both passive and active immunoprophylaxis or immunotherapy. The
     invention further relates to methods for production of the medicaments,
     pharmaceutical compns. containing them and their use in medicine. Also
     forming an aspect of the present invention are ligands, especially monoclonal
     antibodies, which are capable of binding the surface exposed IgE regions
     of the present invention, and their use in medicine as passive
     immunotherapy or in immunoprophylaxis.
     IqE Cepsilon2 epitope mimotope vaccine immunotherapy
ST
IT
     Proteins, specific or class
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (D; epitopes or mimotopes derived from the CE2 domain of IgE
        and antagonists for active and passive immunotherapy of allergic
        disease)
     Immunoglobulins
     RL: BSU (Biological study, unclassified); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (E, Cs2 domain; epitopes or mimotopes derived from the
        Cs2 domain of IgE and antagonists for active and passive
        immunotherapy of allergic disease)
IT
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (PTmAb0005 and PTmAb0011; epitopes or mimotopes derived from the
        CE2 domain of IgE and antagonists for active and passive
        immunotherapy of allergic disease)
IT
     Immunostimulants
        (adjuvants; epitopes or mimotopes derived from the Cs2 domain
        of IgE and antagonists for active and passive immunotherapy of allergic
        disease)
TT
     Purification
        (affinity; epitopes or mimotopes derived from the Cs2 domain of
        IgE and antagonists for active and passive immunotherapy of allergic
        disease)
ΙT
     Diagnosis
        (agents; epitopes or mimotopes derived from the Cs2 domain of
        IgE and antagonists for active and passive immunotherapy of allergic
        disease)
TT
     Antibodies
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (anti-IgE; epitopes or mimotopes derived from the Cs2 domain of
        IgE and antagonists for active and passive immunotherapy of allergic
        disease)
     Drug delivery systems
IT
        (carriers; epitopes or mimotopes derived from the Cs2 domain of
        IqE and antagonists for active and passive immunotherapy of allergic
        disease)
IT
     Adoptive immunotherapy
     Allergy
     Allergy inhibitors
     Blood
     Epitopes
     Immunotherapy
     Protein sequences
     Susceptibility (genetic)
     Vaccines
        (epitopes or mimotopes derived from the Cs2 domain of IgE and
        antagonists for active and passive immunotherapy of allergic disease)
     Fusion proteins (chimeric proteins)
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (epitopes or mimotopes derived from the C\epsilon 2 domain of IgE and
        antagonists for active and passive immunotherapy of allergic disease)
IT
     Antigens
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RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (hepatitis B core; epitopes or mimotopes derived from the \ensuremath{\text{Ce}2}
        domain of IgE and antagonists for active and passive immunotherapy of
        allergic disease)
TT
     Peptides, biological studies
     RL: BSU (Biological study, unclassified); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (mimotope; epitopes or mimotopes derived from the CE2 domain of
        IgE and antagonists for active and passive immunotherapy of allergic
        disease)
     Antibodies
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     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (monoclonal; epitopes or mimotopes derived from the C\epsilon 2 domain
        of IgE and antagonists for active and passive immunotherapy of allergic
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              THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
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RE
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    HCAPLUS
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289886-58-4

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (epitopes or mimotopes derived from the Cs2 domain of IgE and antagonists for active and passive immunotherapy of allergic disease)
 RN 289886-58-4 HCAPLUS
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Absolute stereochemistry.

PAGE 1-B

PAGE 2-A

ANSWER 27 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

2000:102218 HCAPLUS AN

DN 132:245978

Entered STN: 13 Feb 2000 ED

Anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN TI sequence in prostate carcinoma

Livant, Donna L.; Brabec, R. Kaye; Pienta, Kenneth J.; Allen, David L.; ΑU Kurachi, Kotoku; Markwart, Sonja; Upadhyaya, Ameet

Department of Cell and Development Biology, University of Michigan Medical CS School, Ann Arbor, MI, 48109-0616, USA

Cancer Research (2000), 60(2), 309-320 SO CODEN: CNREA8; ISSN: 0008-5472

PB AACR Subscription Office

Journal DT

English LА

CC

1-6 (Pharmacology) Using naturally serum-free SU-ECM basement membranes as invasion AB substrates showed that plasma fibronectin was necessary to stimulate invasion by DU 145 human and metastatic MATLyLu (MLL) rat prostate carcinoma cells. This activity mapped to the PHSRN sequence, which induced invasion through $\alpha5\beta1$ integrin. PHSCN, a competitive inhibitor, blocked both PHSRN- and serum-induced invasion. Acetylated, amidated PHSCN (Ac-PHSCN-NH2) was 30-fold more potent; however, Ac-HSPNC-NH2 was inactive. Rats receiving injections s.c. with 100,000 MLL cells were treated systemically by i.v. injection three times weekly with 1 mg of either Ac-PHSCN-NH2 or Ac-HSPNC-NH2 beginning 24 h later, three times weekly with 1 mg of Ac-PHSCN-NH2 beginning only after surgery to remove large (2 cm) MLL tumors, or were left untreated. MLL tumors grew rapidly in Ac-HSPNC-NH2-treated and in untreated rats. MLL tumor growth in rats treated with Ac-PHSCN-NH2 beginning 1 day after MLL cell injection was reduced by 99.9% during the first 16 days of treatment, although subsequent tumor growth occurred. MLL tumor cryosections immunostained with anti-PECAM-1 showed that Ac-PHSCN-NH2 inhibited neovascularization by 12-fold during this time. Whether initiated after MLL cell injection or only after MLL tumor removal, Ac-PHSCN-NH2 treatment reduced the nos. of MLL lung colonies and micrometastases by 40- to > 100-fold, whereas Ac-HSPNC-NH2 was inactive. Thus, Ac-PHSCN-NH2 may be a potent antitumorigenic and antimetastatic agent for postsurgical use prior to extensive metastasis.

antitumorigenic antimetastatic PHSCN sequence prostate carcinoma; lung ST metastasis inhibitor prostate anticancer AcPHSCNNH2

ΙT Angiogenesis inhibitors

Protein sequences

(anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma)

IT Prostate gland

Prostate gland

(carcinoma, inhibitors; anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma)

ΙT Lung, neoplasm

(inhibitors, metastasis; anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma) IT Antitumor agents (lung, metastasis; anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma) IT Lung, neoplasm (metastasis, inhibitors; anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma) IT Antitumor agents (prostate carcinoma; anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma) IT 262438-43-7 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma) THERE ARE 58 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT RE (1) Akiyama, S; Cancer Metastasis Rev 1995, V14, P173 HCAPLUS (2) Akiyama, S; J Biol Chem 1985, V260, P4492 HCAPLUS (3) Amemiya, S; Dev Growth Differ 1989, V31, P131 (4) Aota, S; J Biol Chem 1994, V269, P24756 HCAPLUS (5) Atherton, E; Solid Phase Peptide Synthesis: A Practical Approach 1989(6) Burnette, W; Anal Biochem 1981, V112, P195 HCAPLUS (7) Carter, H; A Multidisciplinary Analysis of Controversies in the Management of Prostate Cancer 1988, P1 (8) Clark, R; The Molecular and Cellular Biology of Wound Repair 1996, P12 (9) de Souza, P; Br J Cancer 1997, V75, P1593 HCAPLUS (10) Doherty, D; J Clin Invest 1990, V86, P1065 HCAPLUS
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- IT 262438-43-7

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(anti-invasive, antitumorigenic, and antimetastatic activities of the PHSCN sequence in prostate carcinoma)

RN 262438-43-7 HCAPLUS

CN L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

- L24 ANSWER 28 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN
- AN 1999:794362 HCAPLUS
- DN 132:30820
- ED Entered STN: 16 Dec 1999
- TI Anticancer compounds and methods
- IN Livant, Donna L.
- PA Regents of the University of Michigan, USA
- SO U.S., 53 pp., Cont.-in-part of U.S. 5,840,514. CODEN: USXXAM
- DT Patent
- LA English
- IC ICM A61K038-00

ICS A61K038-08

INCL 530330000

CC 1-6 (Pharmacology)

FAN.CNT 4

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os
     MARPAT 132:30820
     The testing of tumor cells, including human tumors capable of metastases,
AB
     in assays employing fibronectin-depleted substrates is described. Ex vivo
     induction of cells, including biopsied human cells, is performed with
     invasion-inducing agents. Addnl., anti-cancer chemotherapeutics are
     described. Specifically, chemotherapeutic agents which have
     anti-metastatic and anti-growth properties are described.
     antitumor peptide metastasis inhibition; prostate cancer metastasis
ST
     antitumor peptide
IT
     Peptides, biological studies
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
        (antitumor peptides and inhibition of metastasis)
IT
     Antitumor agents
        (mammary gland; antitumor peptides and inhibition of metastasis)
IT
     Antitumor agents
        (metastasis; antitumor peptides and inhibition of metastasis)
TT
     Mammary gland
     Mammary gland
     Prostate gland
     Prostate gland
        (neoplasm, inhibitors; antitumor peptides and inhibition of metastasis)
IT
     Antitumor agents
        (prostate gland; antitumor peptides and inhibition of metastasis)
IT
     Fibronectins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
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                                252229-61-1
    252229-59-7
                                              252229-63-3
                                                             252229-65-5
                 252229-60-0
     252229-66-6
                 252229-67-7
                                252229-68-8 252229-69-9
                                                             252229-70-2
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     252229-71-3
                 252229-72-4
     252229-76-8
                 252229-77-9
                                252229-78-0
                                              252229-79-1
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                  252229-82-6
                                252229-83-7
                                               252229-84-8 252229-85-9
     252229-81-5
                                252229-88-2
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     252229-86-0
                  252229-87-1
                 252229-92-8
                                252229-93-9
                                               252229-94-0
                                                             252229-95-1
     252229-91-7
                 252229-97-3
                                252229-98-4
                                               252229-99-5
                                                             252230-00-5
     252229-96-2
     252230-01-6
                 252230-02-7 252230-05-0
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (antitumor peptides and inhibition of metastasis)
     252362-03-1
                  252362-12-2
     RL: PRP (Properties)
       (unclaimed protein sequence; anticancer compds. and methods)
             THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
(1) Anon; WO 9524471 1995 HCAPLUS
(2) Anon; WO 9612823 1996 HCAPLUS
(3) Aversa; US 5576423 1996 HCAPLUS
(4) Bischoff; US 5539085 1996 HCAPLUS
(5) Bohn; US 4424279 1984 HCAPLUS
(6) Bresalier; Cancer Research 1995, V55, P2476 HCAPLUS
(7) Burke; US 5169862 1992 HCAPLUS
(8) Calabresi, P; Goodman and Gilman The Pharmacological Basis of Therapeutics
    8th Edition P1209
(9) Doersen; US 5264358 1993
(10) Douillard; Compendium of Immunology 1981, VII
(11) Eldred; J Med Chem 1994, V37, P3882 HCAPLUS
(12) Gaeta; US 5559103 1996 HCAPLUS
(13) Gartner, T; The Journal of Biological Chemistry V260, P11891 HCAPLUS
(14) Gerlach, J; Cancer Surveys 1986, V5, P25 MEDLINE
(15) Ginsberg; US 5523209 1996 HCAPLUS
(16) Ginsburg; US 5492890 1996 HCAPLUS
(17) Goldie, J; Cancer Research 1984, V44, P3643 HCAPLUS
(18) Hashino; US 5136023 1992 HCAPLUS
(19) Isoai; US 5548062 1996 HCAPLUS
(20) Kitaguchi; US 5436221 1995 HCAPLUS
(21) Kohler, G; European Journal of Immunology 1976, V6, P511 MEDLINE
(22) Kohler, G; Nature 1975, V256, P495 MEDLINE
(23) Ku; J Med Chem 1995, V38, P9 HCAPLUS
(24) Lipman, D; Science 1985, V227, P1435 HCAPLUS
(25) Livant, D; Cancer Research 1995, V55, P5085 HCAPLUS
(26) Lobl; US 5192746 1993 HCAPLUS
(27) Mennen; US 4018653 1977 HCAPLUS
(28) Nicholson, N; Thrombosis Research 1995, V62, P567
(29) Nomizu, M; Cancer Research 1993, V53, P3459 HCAPLUS
(30) Pearson, W; Proc Natl Acad Sci (USA) 1988, V85, P2444 HCAPLUS
(31) Reading, C; Journal of Immunological Methods 1982, V53, P261 MEDLINE
(32) Saiki, I; Cancer Research 1989, V49, P3815 HCAPLUS
(33) Schuurs; US 4016043 1977 HCAPLUS
(34) Shashoua; US 5051448 1991 HCAPLUS
(35) Stone, K; Int J Cancer 1978, V21, P274 MEDLINE
(36) Wenger, R; Blood V73(6), P1498 HCAPLUS
     252229-85-9 252230-05-0
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
```

RE

(Uses)

(antitumor peptides and inhibition of metastasis)

RN 252229-85-9 HCAPLUS

CN L-Asparagine, L-prolyl-L-histidyl-L-seryl-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 252230-05-0 HCAPLUS

CN L-Asparagine, L-prolyl-L-histidyl-L-seryl-L-homocysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L24 ANSWER 29 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1988:72012 HCAPLUS

DN 108:72012

ED Entered STN: 05 Mar 1988

TI Phototransformation pathway of phytochrome

AU Inoue, Y.

CS Fac. Sci., Univ. Tokyo, Tokyo, 113, Japan

SO Springer Proceedings in Physics (1987), 20 (Primary Processes Photobiol.), 70-9

CODEN: SPPPEL; ISSN: 0930-8989

DT Journal; General Review

LA English

CC 11-0 (Plant Biochemistry)

AB A review with 56 refs.

ST review phytochrome phototransformation

IT Photolysis

(of phytochromes, pathway of)

IT Phytochromes

RL: BIOL (Biological study)

(phototransformation pathway of)
IT 73508-81-3
 RL: BIOL (Biological study)
 (phototransformation pathway of)
IT 73508-81-3
 RL: BIOL (Biological study)
 (phototransformation pathway of)
RN 73508-81-3 HCAPLUS
CN L-Tyrosine, N-[N2-[N-[N-[S-[1-[8,12-bis(2-carboxyethyl)-18-ethenyl-1,2,3,19,22,24-hexahydro-2,7,13,17-tetramethyl-1,19-dioxo-21H-bilin-3-yl]ethyl]-N-[N-[N-[1-[N-(N2-L-leucyl-L-arginyl)-L-alanyl]-L-prolyl]-L-histidyl]-L-seryl]-L-cysteinyl]-L-histidyl]-L-glutaminyl]-, [2R,3R(R)]- (9CI) (CA INDEX NAME)

— он

i-Bu

PAGE 2-A

 ${\tt HO_2C-CH_2-}$

PAGE 2-B

$$CH$$
 CH
 CH
 CH_2
 CH

PAGE 3-B

н₂с== Сн

L24 ANSWER 30 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN 1987:405411 HCAPLUS ANDN107:5411 ED Entered STN: 11 Jul 1987 Molecular interactions in the "GAT" idiotypic network: an approach using TI synthetic peptides Mazza, G.; Guigou, V.; Moinier, D.; Corbet, S.; Ollier, P.; Fougereau, M. ΑU Cent. Immunol., INSERM-CNRS Marseille-Luminy, Marseille, 13288, Fr. CS Annales de l'Institut Pasteur/Immunology (1987), 138(1), 3-17 SO CODEN: AIPIEP; ISSN: 0769-2625 DTJournal LA English CC 15-2 (Immunochemistry)

In order to approach some of the dominant epitopes which are recognized in the GAT (Glu60Ala30Tyr10)n random terpolymer, a variety of peptides containing 7-14 residues were synthesized using glutamic acid and tyrosine as building blocks, and thus were able to mimic determinants common to GAT and GT (Glu50Tyr50)n. One decapeptide and one dodecapeptide inhibited GT-mAb1 (or mAb1') binding to the same extent as GAT. Antibodies were also raised against synthetic peptides which reproduced the sequence of the 6 complementarity-determining regions of the germline anti-GAT Ab1 antibody. Antibodies were obtained against all peptides except L1, and were shown to recognize the native Ab1-Fab. Surprisingly, some of these antibodies also recognized GAT, i.e. anti-L2, anti-H2, and anti-H3, an observation which speaks in favor of a triggering of the idiotypic network at the Ab3 level. Finally, a monoclonal antibody derived from an immunization with an Ab2-D region synthetic peptide was found to be of the Ag+Id- type. Sequence data indicate that the light chain at least is completely different from that of the Ab1/Ab1', which uses only a very precise pair of V germline genes.

GAT peptide antibody idiotype ST Antibodies IT RL: PREP (Preparation) (GAT idiotypic network of, preparation of peptides for study of) Peptides, preparation IT RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study); PREP (Preparation) (preparation and antigenicity of, for GAT idiotypic network study) TT 31325-29-8, GAT RL: BIOL (Biological study) (antibody idiotypic network system involving, mol. interactions in, preparation of peptides for study of) 108526-52-9P 108526-53-0P 108526-54-1P IT 108526-51-8P 108526-55-2P 108543-65-3P RL: PREP (Preparation) (preparation and antibodies binding to, in GAT idiotypic network studies) IT 31325-39-0P RL: PREP (Preparation) (preparation and antigenicity of peptides like, in GAT idiotypic network studies) 108526-59-6P 108526-60-9P 108526-56-3P 108526-57-4P 108526-58-5P IT 108526-61-0P RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study); PREP (Preparation) (preparation and antigenicity of, in GAT idiotypic network studies) IT 108526-54-1P RL: PREP (Preparation) (preparation and antibodies binding to, in GAT idiotypic network studies) RN 108526-54-1 HCAPLUS L-Cysteine, N-[N-[N-[1-[N-[N-[N-[N-(N2-L-phenylalanyl-L-glutaminyl)glycyl]-CN L-threonyl]-L-histidyl]-L-valyl]-L-prolyl]-L-histidyl]-L-threonyl]- (9CI)

Absolute stereochemistry.

(CA INDEX NAME)

PAGE 1-A

PAGE 2-A

L24 ANSWER 31 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

```
ΑN
     1986:604942 HCAPLUS
     105:204942
     Entered STN: 13 Dec 1986
ED
     Site-directed antisera to the chromophore binding site of phytochrome:
ΤI
     characterization and cross-reactivity
     Mercurio, Frank M.; Houghten, Richard A.; Lagarias, J. Clark
ΑIJ
     Dep. Biochem. Biophys., Univ. California, Davis, CA, 95616, USA
CS
     Archives of Biochemistry and Biophysics (1986), 248(1), 35-42
so
     CODEN: ABBIA4; ISSN: 0003-9861
DT
     Journal
LΑ
     English
CC
     6-3 (General Biochemistry)
     Section cross-reference(s): 11, 15
     The preparation and characterization of antisera to a synthetic undecapeptide
AB
     which encompasses the chromophore binding site on oat phytochrome are
     described. By direct enzyme-linked immunosorbent assay, this antiserum
     recognizes bilin-linked peptides in proteolytic digestion mixts. of
     phytochrome from several plant species including the monocot species, oat
     and maize, and dicot species, pea. Competitive enzyme-linked
     immunosorbent assays were used to establish the specificity of the
     site-directed antisera for those peptides homologous with the synthetic
     undecapeptide which are present in complex mixts. Apparently, the local
     environment of the phytochromobilin prosthetic group is structurally
     conserved.
ST
     phytochrome chromophore binding site
IT
     Phytochromes
     RL: BIOL (Biological study)
        (chromophore binding site of, characterization of)
IT
     Oat
     Pea
        (phytochrome of, chromophore binding site of, characterization of)
IT
     Evolution
```

(conservative, of phytochrome chromophore binding site)

- IT Antibodies
 - RL: BIOL (Biological study)

(monoclonal, to chromophore binding site of phytochrome)

- IT 73508-81-3 103516-77-4
 - RL: RCT (Reactant); RACT (Reactant or reagent)

(reaction of, with antiserum, characterization of chromophore binding site of phytochrome in relation to) ${\bf r}$

- IT 73508-81-3 103516-77-4
 - RL: RCT (Reactant); RACT (Reactant or reagent)

(reaction of, with antiserum, characterization of chromophore binding site of phytochrome in relation to)

- RN 73508-81-3 HCAPLUS
- CN L-Tyrosine, N-[N2-[N-[S-[1-[8,12-bis(2-carboxyethyl)-18-ethenyl-1,2,3,19;22,24-hexahydro-2,7,13,17-tetramethyl-1,19-dioxo-21H-bilin-3-yl]ethyl]-N-[N-[N-[1-[N-(N2-L-leucyl-L-arginyl)-L-alanyl]-L-prolyl]-L-histidyl]-L-seryl]-L-cysteinyl]-L-histidyl]-L-leucyl]-L-glutaminyl]-,

[2R,3R(R)]- (9CI) (CA INDEX NAME)

— он

i-Bu

PAGE 2-A

но₂с-сн₂-

$$CH$$
 CH
 CH
 $CH_2 - CH_2 - CO_2H$
 Me
 CH
 Me
 NH
 NH

ĊN prolyl]-L-histidyl]-L-seryl]-L-cysteinyl]-L-histidyl]-L-leucyl]-L-glutaminyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L24 ANSWER 32 OF 32 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 1980:463852 HCAPLUS

DN 93:63852

ED Entered STN: 12 May 1984

TI Chromopeptides from phytochrome. The structure and linkage of the PR form of the phytochrome chromophore

AU Lagarias, J. Clark; Rapoport, Henry

CS Dep. Chem., Univ. California, Berkeley, CA, 94720, USA

SO Journal of the American Chemical Society (1980), 102(14), 4821-8

CODEN: JACSAT; ISSN: 0002-7863

DT Journal

LA English

CC 6-3 (General Biochemistry)

GΙ

The isolation and chromatog, purification of chromophore-containing peptides from the PR form of phytochrome treated with pepsin and thermolysin are described. From the amino acid sequence and 1H NMR spectral anal. of phytochromobiliundecapeptide (I), the structure of the PR phytochrome chromophore and the nature of the thioether linkage joining pigment to peptide were established. Confirmatory evidence was obtained from similar anal. of phytochromobilioctapeptide. The implications of this structural assignment with respect to the mechanism of the PR to PFR phototransformation are considered.

```
phytochrome PR chromophore structure
ST
IT
     Peptides, properties
     RL: PRP (Properties)
        (amino acid sequence of, of phytochrome PR chromopeptide)
IT
     Nuclear magnetic resonance
     Ultraviolet and visible spectra
         (of phytochrome PR chromopeptide)
IT
        (phytochrome PR of, chromophore structure of)
     Phytochromes
IT
        (PR, chromophore of, structure and photochem. transformation of)
IT
     73508-81-3
     RL: BIOL (Biological study)
        (of phytochrome PR)
     73508-81-3
IT
     RL: BIOL (Biological study)
         (of phytochrome PR)
     73508-81-3 HCAPLUS
L-Tyrosine, N-[N2-[N-[N-[S-[1-[8,12-bis(2-carboxyethyl)-18-ethenyl-
RN
CN
     1,2,3,19,22,24-hexahydro-2,7,13,17-tetramethyl-1,19-dioxo-21H-bilin-3-
     yl]ethyl]-N-[N-[N-[N-[N-(N2-L-leucyl-L-arginyl)-L-alanyl]-L-prolyl]-L-
     histidyl]-L-seryl]-L-cysteinyl]-L-histidyl]-L-leucyl]-L-glutaminyl]-,
     [2R,3R(R)]- (9CI) (CA INDEX NAME)
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— он

i-Bu

PAGE 2-A

но₂с-сн₂-

PAGE 2-B

$$CH$$
 CH
 CH_2
 CH_2

PAGE 3-B

Н2С==СН

=> b uspat
'USPAT' IS AN AMBIGUOUS FILE OR CLUSTER NAME
USPATALL - US Patent Full Text Cluster
USPAT2 - U.S. Patents Latest Publications from 2001 - present
USPATFULL - U.S. Patents Original Publications from 1971 - present
ENTER FILE OR CLUSTER NAME (IGNORE):end

=> b uspatall FILE 'USPATFULL' ENTERED AT 08:57:07 ON 03 AUG 2005 CA INDEXING COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'USPAT2' ENTERED AT 08:57:07 ON 03 AUG 2005 CA INDEXING COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

=> d bib abs fhitstr 135 tot

L35 ANSWER 1 OF 2 USPATFULL on STN 2005:24260 USPATFULL ΑN Peptides which target tumor and endothelial cells, compositions and uses ΤI thereof Ternansky, Robert J., San Diego, CA, UNITED STATES IN Allan, Amy L., Encinitas, CA, UNITED STATES Gladstone, Patricia L., San Diego, CA, UNITED STATES Yoon, Won Hyung, San Diego, CA, UNITED STATES Parry, Graham, San Diego, CA, UNITED STATES Donate, Fernando, San Diego, CA, UNITED STATES Mazar, Andrew, San Diego, CA, UNITED STATES US 2005020810 20050127 PΙ A1 US 2003-722843 20031125 (10) ΑI Α1

PRAI US 2002-429174P 20021125 (60)

US 2003-475539P 20030602 (60)

DT Utility

FS APPLICATION

LREP Sunil K: Singh, Dorsey & Whitney LLP, Intellectual Property Department,

Four Embarcadero Center, Suite 3400, San Francisco, CA, 94111-4187

CLMN Number of Claims: 74

ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 3884

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention relates generally to peptide analogs of Ac-PHSCN--NH.sub.2 which target tumor and endothelial cells and have anti-tumor, anti-angiogenic and anti-metasastic activity, methods of making these peptides, compositions thereof and methods of using these peptides and pharmaceutical compositions thereof to treat, prevent and detect diseases characterized by tumor growth, metastasis and angiogenesis. The peptide analogs may serve, inter alia, as carriers of radioactivity, PET-active compounds, toxins, fluorescent molecules and PEG molecules.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 701200-82-0P

(peptide inhibitors of angiogenesis, cell migration, cell invasion and cell proliferation, preparation, and compns. and therapeutic uses)

RN 701200-82-0 USPATFULL

CN L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-S-[(4-methylphenyl)methyl]-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L35 ANSWER 2 OF 2 USPATFULL on STN

AN 2004:209805 USPATFULL

TI Peptides which inhibit angiogenesis, cell migration, cell invasion and cell proliferation, compositions and uses thereof

IN Allan, Amy L., Encinitas, CA, UNITED STATES

Donate, Fernando, San Diego, CA, UNITED STATES Hopkins, Stephanie A., Poway, CA, UNITED STATES

Gladstone, Patricia L., San Diego, CA, UNITED STATES

Mazar, Andrew, San Diego, CA, UNITED STATES

O'Hare, Sean M., San Diego, CA, UNITED STATES

Parry, Graham, San Diego, CA, UNITED STATES

Plunkett, Marian, San Diego, CA, UNITED STATES

Ternansky, Robert J., San Diego, CA, UNITED STATES Yoon, Won Hyung, San Diego, CA, UNITED STATES

PI US 2004162239 A1 20040819 AI US 2003-723144 A1 20031125 (10) PRAI US 2002-429174P 20021125 (60) US 2003-475539P 20030602 (60) DT Utility

FS APPLICATION

LREP COOLEY GODWARD, LLP, 3000 EL CAMINO REAL, 5 PALO ALTO SQUARE, PALO ALTO,

CA, 94306

CLMN Number of Claims: 65 ECL Exemplary Claim: 1 DRWN 5 Drawing Page(s)

LN.CNT 3373

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention relates generally to peptides, which inhibit angiogenesis, cell migration, cell invasion and cell proliferation, methods of making peptides, which inhibit angiogenesis, cell migration, cell invasion and cell proliferation, pharmaceutical compositions of these peptides and methods of using these peptides and pharmaceutical compositions of these peptides to treat diseases associated with aberrant vascularization.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 701200-82-0P

(peptide inhibitors of angiogenesis, cell migration, cell invasion and cell proliferation, preparation, and compns. and therapeutic uses)

RN 701200-82-0 USPATFULL

CN L-Aspartamide, 1-acetyl-L-prolyl-L-histidyl-L-seryl-S-[(4-methylphenyl)methyl]-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

=> d bib abs fhitstr hitrn 137 tot

L37 ANSWER 1 OF 9 USPATFULL on STN

AN 2005:117696 USPATFULL

TI KDR and VEGF/KDR binding peptides and their use in diagnosis and therapy

IN Sato, Aaron K., Somerville, MA, UNITED STATES
Sexton, Daniel J., Melrose, MA, UNITED STATES
Dransfield, Daniel T., Hanson, MA, UNITED STATES
Ladner, Robert C., Ijamsville, MD, UNITED STATES
Arbogast, Christophe, Viuz - En - Sallaz, FRANCE
Bussat, Philippe, Feigers, FRANCE
Fan, Hong, Plainsboro, NJ, UNITED STATES

Khurana, Sudha, Plainsboro, NJ, UNITED STATES

```
Linder, Karen E., Kingston, NJ, UNITED STATES
       Marinelli, Edmund R., Lawrenceville, NJ, UNITED STATES
       Nanjappan, Palaniappa, Dayton, NJ, UNITED STATES
       Nunn, Adrian D., Lamberville, NJ, UNITED STATES
       Pillai, Radhakrishna, Cranbury, NJ, UNITED STATES
       Pochon, Sibylle, Troinex, SWITZERLAND
       Ramalingam, Kondareddiar, Dayton, NJ, UNITED STATES
       Shrivastava, Ajay, Plainsboro, NJ, UNITED STATES
       Song, Bo, Princeton, NJ, UNITED STATES
       Swenson, Rolf E., Princeton, NJ, UNITED STATES
       Von Wronski, Mathew A., Moorestown, NJ, UNITED STATES
       Dyax Corporation, Cambridge, MA, UNITED STATES (U.S. corporation)
PA
       Bracco International B. V. (U.S. corporation)
                               20050512
PΙ
       US 2005100963
                         A1
                               20030911 (10)
AΙ
       US 2003-661156
                         A1
       Continuation-in-part of Ser. No. US 2003-382082, filed on 3 Mar 2003,
RLT
       ABANDONED Continuation-in-part of Ser. No. WO 2003-US6731, filed on 3
       Mar 2003, PENDING
       US 2002-360851P
                           20020301 (60)
                                                                     <--
PRAI
       US 2003-440411P
                           20030115 (60)
DT
       Utility
FS
       APPLICATION
       HAMILTON, BROOK, SMITH & REYNOLDS, P.C., 530 VIRGINIA ROAD, P.O. BOX
LREP
       9133, CONCORD, MA, 01742-9133, US
CLMN
       Number of Claims: 199
       Exemplary Claim: 1
ECL
DRWN
       118 Drawing Page(s)
LN.CNT 14344
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention provides polypeptides, peptide dimer, and
AΒ
       multimeric complexes comprising at least one binding moiety for KDR or
       VEGF/KDR complex, which have a variety of uses wherever treating,
       detecting, isolating or localizing angiogenesis is advantageous.
       Particularly disclosed are synthetic, isolated polypeptides capable of
       binding KDR or VEGF/KDR complex with high affinity (e.g., having a
       K.sub.D<1~\mu M), and dimer and multimeric constructs comprising these
```

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 599210-01-2

(amino acid sequence; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

RN 599210-01-2 USPATFULL

polypeptides.

CN L-Glutamic acid, glycyl-L-histidyl-L-prolyl-L-cysteinyl-L-lysylglycyl-L-methionyl-L-leucyl-L-prolyl-L-histidyl-L-threonyl-L-cysteinyl-L-tryptophyl-L-tyrosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

IT 599210-01-2 851724-81-7

(amino acid sequence; sequences of KDR and VEGF/KDR binding peptides, peptide dimers, and multimeric complexes and their use in diagnosis and therapy)

L37 ANSWER 2 OF 9 USPATFULL on STN

2005:68857 USPATFULL ΑN

Binding phenol oxidizing enzyme-peptide complexes ΤI

IN Wolfgang, Aehle, Delfgauw, NETHERLANDS

Baldwin, Toby M., Palo Alto, CA, UNITED STATES

Van Gastel, Franciscus J.C., Union City, CA, UNITED STATES

Janssen, Giselle G., San Carlos, CA, UNITED STATES

Murray, Christopher J., Soquel, CA, UNITED STATES

Wang, Huaming, Fremont, CA, UNITED STATES

Winetzky, Deborah S., Foster City, CA, UNITED STATES

20050317 PΤ US 2005058996 Α1

20020903 (10) ΑI

US 2002-235043 Α1

DTUtility

APPLICATION FS

H. THOMAS ANDERTON, JR., Genencor International, Inc., 925 Page Mill LREP Road, Palo Alto, CA, 94034-1013

Number of Claims: 30 CLMN

ECL Exemplary Claim: 1

10 Drawing Page(s) DRWN

LN.CNT 3422

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present application relates to peptides which bind to a target AB stain, phenol oxidizing enzyme-binding peptide complexes wherein the binding peptide is attached to the C-terminus of the phenol oxidizing enzyme or is inserted or substituted into the phenol oxidizing enzyme. In a preferred embodiment the phenol oxidizing enzyme is a laccase specifically Stachybotrys oxidase B and variants thereof. The invention provides expression vectors comprising the phenol oxidizing enzyme-binding peptide complex as well as host cells comprising the vectors.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

502166-83-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme

(amino acid sequence; peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes for creation of detergent)

RN 502166-83-8 USPATFULL

L-Methionine, L-isoleucyl-L-asparaginyl-L-threonyl-L-prolyl-L-histidyl-L-CN seryl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

TT 502166-83-8DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-78-4DP, cysteine derivs. and fusion product with phenol oxidizing enzyme 502167-93-3DP, cysteine derivs. and fusion product with phenol oxidizing enzyme (amino acid sequence; peptides able to bind carotenoid compds., their sequences and use in construction of fusion proteins with Stachybotrys phenol oxidizing enzymes for creation of detergent)

```
ANSWER 3 OF 9 USPATFULL on STN
L37
       2005:23322 USPATFULL
AN
       52 Human secreted proteins
TI
       Ni, Jian, Germantown, MD, UNITED STATES
IN
       Baker, Kevin P., Darnestown, MD, UNITED STATES
       Birse, Charles E., North Potomac, MD, UNITED STATES
       Fiscella, Michele, Bethesda, MD, UNITED STATES
       Komatsoulis, George A., Silver Spring, MD, UNITED STATES
       Rosen, Craig A., Laytonsville, MD, UNITED STATES
       Soppet, Daniel R., Centreville, VA, UNITED STATES
       Young, Paul E., Gaithersburg, MD, UNITED STATES
       Ebner, Reinhard, Gaithersburg, MD, UNITED STATES
       Duan, D. Roxanne, Gaithersburg, MD, UNITED STATES
       Olsen, Henrik S., Gaithersburg, MD, UNITED STATES
       La Fleur, David W., Washington, DC, UNITED STATES
       Moore, Paul A., North Bethesda, MD, UNITED STATES
       Shi, Yanggu, Gaithersburg, MD, UNITED STATES
       Wei, Ping, Agoura Hills, CA, UNITED STATES
       Florence, Kimberly A., Rockville, MD, UNITED STATES
       Human Genome Sciences, Inc., Rockville, MD, UNITED STATES, 20850 (U.S.
PA
       corporation)
                               20050127
       US 2005019866
PΙ
                          A1
       US 2004-883936
                          A1
                               20040706 (10)
AΙ
       Continuation of Ser. No. US 2001-789561, filed on 22 Feb 2001, PENDING
RLI
       Continuation-in-part of Ser. No. WO 2000-US24008, filed on 31 Aug 2000,
       PENDING
       US 1999-152317P
                           19990903 (60)
                                                                     < - -
PRAI
                           19990903 (60)
       US 1999-152315P
DT
       Utility
FS
       APPLICATION
       HUMAN GENOME SCIENCES INC, INTELLECTUAL PROPERTY DEPT., 14200 SHADY
LREP
       GROVE ROAD, ROCKVILLE, MD, 20850
CLMN
       Number of Claims: 24
ECL
       Exemplary Claim: 1
       No Drawings
DRWN
LN.CNT 24739
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
```

The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 328529-77-7

(unclaimed sequence; cloning and cDNA and deduced amino acid sequences of 52 human secreted proteins)

RN 328529-77-7 USPATFULL

CN Glycine, L-lysyl-L-tryptophylglycyl-L-cysteinyl-L-phenylalanyl-L-cysteinylL-lysylglycyl-L-seryl-L-seryl-L-phenylalanyl-L-threonyl-L-prolyl-Lhistidyl-L-seryl-L-cysteinyl-L-prolyl-L-prolyl-L-α-glutamyl-Lalanyl-L-prolyl-L-leucyl-L-phenylalanyl-L-prolyl-L-alanyl-L-valyl-Lleucyl-L-leucyl-L-seryl-L-threonyl-L-leucyl- (9CI) (CA INDEX
NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

PAGE 1-C

PAGE 1-E

_CO2H

PAGE 2-D

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IT
     328529-77-7
        (unclaimed sequence; cloning and cDNA and deduced amino acid sequences
        of 52 human secreted proteins)
    ANSWER 4 OF 9 USPATFULL on STN
       2004:2568 USPATFULL
ΑN
TI
       50 human secreted proteins
       Moore, Paul A., Germantown, MD, UNITED STATES
IN
       Ruben, Steven M., Olney, MD, UNITED STATES
       LaFleur, David W., Washington, DC, UNITED STATES
       Shi, Yanggu, Gaithersburg, MD, UNITED STATES
       Rosen, Craig A., Laytonsville, MD, UNITED STATES
       Olsen, Henrik S., Gaithersburg, MD, UNITED STATES
       Ebner, Reinhard, Gaithersburg, MD, UNITED STATES
       Brewer, Laurie A., St. Paul, MN, UNITED STATES
       Human Genome Sciences, Inc., Rockville, MD (U.S. corporation)
PA
       US 2004002591
                         A1 20040101
PΙ
AΤ
       US 2002-47021
                         A1
                               20020117 (10)
       Continuation-in-part of Ser. No. US 2000-722329, filed on 28 Nov 2000,
RLT
       PENDING Continuation of Ser. No. US 1999-262109, filed on 4 Mar 1999,
       ABANDONED Continuation-in-part of Ser. No. WO 1998-US18360, filed on 3
       Sep 1998, PENDING
PRAT
       US 2001-262066P
                           20010118 (60)
                                                                      <--
                                                                      <---
       US 1997-57626P
                           19970905 (60)
                                                                      <---
       US 1997-57663P
                           19970905 (60)
                           19970905 (60)
       US 1997-57669P
                                                                      <--
       US 1997-58666P
                           19970912 (60)
       US 1997-58667P
                           19970912 (60)
                                                                      <--
                           19970912 (60)
                                                                      < - -
       US 1997-58973P
                           19970912 (60)
                                                                      <---
       US 1997-58974P
       US 1998-90112P
                                                                      <--
                           19980622 (60)
DT
       Utility
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Exemplary Claim: 1 DRWN 2 Drawing Page(s) LN.CNT 33379

Number of Claims: 23

CAS INDEXING IS AVAILABLE FOR THIS PATENT. The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells,

antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

443339-97-7

(unclaimed sequence; cloning and cDNA and deduced amino acid sequences of 50 human secreted proteins)

443339-97-7 USPATFULL RN

APPLICATION

FS

LREP CLMN

ECL

L-Valine, L-alanyl-L-leucyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-CN $leucyl-L-\alpha-glutamyl-L-arginyl-L-alanyl-L-lysyl-L-\alpha-glutamyl-$ L-isoleucyl-L-lysyl-L-isoleucyl-L-lysyl-L-leucylglycyl-L-isoleucyl-Lleucyl-L-leucyl-L-glutaminyl-L-lysyl-L-prolyl-L-α-aspartyl-L-seryl-L-valylglycyl-L-α-aspartyl-L-leucyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

$$H_{2N}$$
 CH_{2}
 H_{2N}
 CH_{2}
 H_{2N}
 H

PAGE 1-C

PAGE 1-D

PAGE 1-E

IT 443339-97-7

L37

(unclaimed sequence; cloning and cDNA and deduced amino acid sequences of 50 human secreted proteins)

AN 2003:173232 USPATFULL
TI B7-like polynucleotides, polypeptides, and antibodies
IN Ruben, Steven M., Olney, MD, UNITED STATES
Chen, Lieping, Rochester, MN, UNITED STATES
Baker, Kevin P., Darnestown, MD, UNITED STATES

Ni, Jian, Germantown, MD, UNITED STATES
PA Human Genome Sciences, Inc., Rockville, MD, UNITED STATES, 20850 (U.S.

corporation)

PI US 2003119076 A1 20030626

ANSWER 5 OF 9 USPATFULL on STN

AI US 2002-141953 A1 20020510 (10) <--

RLI Continuation of Ser. No. US 2001-790622, filed on 23 Feb 2001, PENDING Continuation-in-part of Ser. No. WO 2000-US23792, filed on 30 Aug 2000, PENDING

PRAI US 1999-152317P 19990903 (60) <--US 2000-200346P 20000428 (60) <--

DT Utility

FS APPLICATION

LREP HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850

CLMN Number of Claims: 22 ECL Exemplary Claim: 1 DRWN 15 Drawing Page(s)

LN.CNT 12418

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention relates to novel human B7-like polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human B7-like polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human B7-like polypeptides.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 328529-77-7

(unclaimed sequence; cloning and cDNA and deduced amino acid sequences of 52 human secreted proteins)

RN 328529-77-7 USPATFULL

CN Glycine, L-lysyl-L-tryptophylglycyl-L-cysteinyl-L-phenylalanyl-L-cysteinyl-L-lysylglycyl-L-seryl-L-seryl-L-phenylalanyl-L-threonyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-prolyl-L-prolyl-L-α-glutamyl-L-alanyl-L-prolyl-L-leucyl-L-phenylalanyl-L-prolyl-L-alanyl-L-valyl-L-leucyl-L-leucyl-L-valyl-L-seryl-L-threonyl-L-leucyl- (9CI) (CA INDEX

Absolute stereochemistry.

PAGE 1-A

PAGE 1-B

PAGE 1-C

PAGE 1-E

__CO2H

PAGE 2-D

328529-77-7 IT

(unclaimed sequence; cloning and cDNA and deduced amino acid sequences of 52 human secreted proteins)

ANSWER 6 OF 9 USPATFULL on STN L37

2003:146720 USPATFULL AN

Binding phenol oxidizing enzyme-peptide complexes TI

Aehle, Wolfgang, Delfgauw, NETHERLANDS IN

Baldwin, Toby M., Palo Alto, CA, UNITED STATES

van Gastel, Franciscus J. C., Union City, CA, UNITED STATES

Janssen, Giselle G., San Carlos, CA, UNITED STATES Murray, Christopher J., Soquel, CA, UNITED STATES Wang, Huaming, Fremont, CA, UNITED STATES

Winetzky, Deborah S., Foster City, CA, UNITED STATES

20030529 A1 ΡI US 2003100467

US 2001-954385 A1 20010912 (9) AΙ

Utility DT

APPLICATION FS

GENENCOR INTERNATIONAL, INC., 925 PAGE MILL ROAD, PALO ALTO, CA, 94304 LREP

Number of Claims: 39 CLMN

Exemplary Claim: 1 ECL

,11 Drawing Page(s) DRWN

LN.CNT 1461

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present application relates to peptides which bind to a target stain, phenol oxidizing enzyme-binding peptide complexes wherein the binding peptide is attached to the C-terminus of the phenol oxidizing enzyme or is inserted or substituted into the phenol oxidizing enzyme. In a preferred embodiment the phenol oxidizing enzyme is a laccase specifically Stachybotrys oxidase B and variants thereof. The invention provides expression vectors comprising the phenol oxidizing enzyme-binding peptide complex as well as host cells comprising the vectors.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

502169-28-0

(protein motif; selection and production of laccase- and other phenol oxidizing enzyme-binding peptide complexes for detergents and carotenoid stain removal)

502169-28-0 USPATFULL

L-Methionine, L-prolyl-L-histidyl-L-seryl- (9CI) (CA INDEX NAME) CN

Absolute stereochemistry.

502169-28-0 IT

(protein motif; selection and production of laccase- and other phenol

Search done by Noble Jarrell

oxidizing enzyme-binding peptide complexes for detergents and carotenoid stain removal)

IT 502166-83-8P 502167-78-4P 502167-93-3P

(selection and production of laccase- and other phenol oxidizing enzyme-binding peptide complexes for detergents and carotenoid stain removal)

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L37 ANSWER 7 OF 9 USPATFULL on STN
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AN 2002:307551 USPATFULL

TI Mannosidase structures

IN Rose, David Richard, Markham, CANADA Kuntz, Douglas Arthur, Toronto, CANADA

Van Den Elsen, Jean Maria Hubertus, Bath, UNITED KINGDOM

PI US 2002172670 A1 20021121 <--AI US 2001-960226 A1 20010921 (9) <--PRAI US 2001-263458P 20010123 (60) <--US 2000-234879P 20000922 (60) <---

DT Utility

FS APPLICATION

LREP MERCHANT & GOULD PC, P.O. BOX 2903, MINNEAPOLIS, MN, 55402-0903

CLMN Number of Claims: 47 ECL Exemplary Claim: 1 DRWN 20 Drawing Page(s)

LN.CNT 40572

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention relates to a crystal comprising a mannosidase II ligand-binding domain. In particular the present invention relates to a crystal comprising mannosidase II (with and without swainsonine), and its use to generate models for elucidating the structure of other polypeptides and for better identifying ligands capable of modulating mannosidase II activity.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 476173-05-4

(protein motif; crystal structure of Drosophila α -mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

RN 476173-05-4 USPATFULL

Absolute stereochemistry.

PAGE 1-C

$$H_2N$$
 S
 H
 S
 H

PAGE 2-A

$$\begin{array}{c} & & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & &$$

IT 476173-05-4

(protein motif; crystal structure of Drosophila α-mannosidase II and swainsonine complexes, its use for identifying mannosidase II activity-modulating ligands and therapeutic applications)

ANSWER 8 OF 9 USPATFULL on STN L37

2002:126306 USPATFULL ΑN

ΤI 52 human secreted proteins

Ni, Jian, Germantown, MD, UNITED STATES IN Baker, Kevin P., Darnestown, MD, UNITED STATES Birse, Charles E., North Potomac, MD, UNITED STATES Fiscella, Michele, Bethesda, MD, UNITED STATES Komatsoulis, George A., Silver Spring, MD, UNITED STATES Rosen, Craig A., Laytonsville, MD, UNITED STATES Soppet, Daniel R., Centreville, VA, UNITED STATES Young, Paul E., Gaithersburg, MD, UNITED STATES Ebner, Reinhard, Gaithersburg, MD, UNITED STATES Duan, D. Roxanne, Bethesda, MD, UNITED STATES Olsen, Henrik S., Gaithersburg, MD, UNITED STATES LaFleur, David W., Washington, DC, UNITED STATES Moore, Paul A., Germantown, MD, UNITED STATES

Shi, Yanggu, Gaithersburg, MD, UNITED STATES

Wei, Ping, Brookeville, MD, UNITED STATES Florence, Kimberly A., Rockville, MD, UNITED STATES

PI US 2002064818 A1 20020530 <--AI US 2001-789561 A1 20010222 (9) <--

RLI Continuation-in-part of Ser. No. WO 2000-US24008, filed on 31 Aug 2000,

UNKNOWN

PRAI US 1999-152317P 19990903 (60) <--US 1999-152315P 19990903 (60) <--

DT Utility

FS APPLICATION

LREP HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850

CLMN Number of Claims: 23 ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 24623

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 328529-77-7

(unclaimed sequence; cloning and cDNA and deduced amino acid sequences of 52 human secreted proteins)

RN 328529-77-7 USPATFULL

CN Glycine, L-lysyl-L-tryptophylglycyl-L-cysteinyl-L-phenylalanyl-L-cysteinyl-L-lysylglycyl-L-seryl-L-seryl-L-phenylalanyl-L-threonyl-L-prolyl-L-histidyl-L-seryl-L-cysteinyl-L-prolyl-L-prolyl-L-α-glutamyl-L-alanyl-L-prolyl-L-leucyl-L-phenylalanyl-L-prolyl-L-alanyl-L-valyl-L-leucyl-L-leucyl-L-benyl-L-threonyl-L-leucyl-L-leucyl-L-valyl-L-seryl-L-threonyl-L-leucyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

$$H_2N$$
 S (CH_2) A NH_2 SH SH NH_2 N

PAGE 1-C

PAGE 1-E

-CO2H

PAGE 2-D

IT 328529-77-7

(unclaimed sequence; cloning and cDNA and deduced amino acid sequences of 52 human secreted proteins)

ANSWER 9 OF 9 USPATFULL on STN 1999:163820 USPATFULL L37 ANAnticancer compounds and methods ΤI Livant, Donna L., Ann Arbor, MI, United States IN The Regents of the University of Michigan, Ann Arbor, MI, United States PA (U.S. corporation) ΡI US 6001965 19991214 US 1997-915189 19970820 (8) <--ΑI Continuation-in-part of Ser. No. US 1996-754322, filed on 21 Nov 1996, RLI now patented, Pat. No. US 5840514, issued on 24 Nov 1998 DT Utility Granted FS Primary Examiner: Woodward, Michael P.; Assistant Examiner: Borin, EXNAM Michael LREP Medlen & Carroll, LLP Number of Claims: 5 CLMN ECL Exemplary Claim: 1

DRWN 16 Drawing Figure(s); 15 Drawing Page(s)

LN.CNT 2294

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The testing of tumor cells, including human tumors capable of metastases, in assays employing fibronectin-depleted substrates is described. Ex vivo induction of cells, including biopsied human cells, is performed with invasion-inducing agents. Additionally, anti-cancer chemotherapeutics are described. Specifically, chemotherapeutic agents which have anti-metastatic and anti-growth properties are described.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 252229-85-9

(antitumor peptides and inhibition of metastasis)

RN 252229-85-9 USPATFULL

CN L-Asparagine, L-prolyl-L-histidyl-L-seryl-L-cysteinyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 252229-85-9 252230-05-0 (antitumor peptides and inhibition of metastasis)

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Search done by Noble Jarrell